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620 NEWPORT CENTER DRIVE

SIXTEENTH FLOOR

NEWPORT BEACH, CALIFORNIA 92660-8016

(949) 760-0404

FAX (949) 760-9502

INTERNET: WWW.KMOB.COM

LOUIS J. KNOBBE*
 DON W. MARTENS
 GORDON H. OLSON†
 JAMES D. BEAR
 DALE C. OLSON*
 RANDY JUNKER
 MICHAEL NIEMAN
 CHRISTOPHER PETERSON
 ROSE R. ROSE*
 MICHAEL SNIAK
 RAELESEN
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 MARGUERITE L. GUNN
 STEPHEN C. JENSEN
 VITO A. CANUSO

Assistant Commissioner for Patents

Washington, D.C. 20231

ANN A. BYUN
 ROBERT F. GAZDZINSKI
 FRED C. HERNANDEZ
 STACEY R. HALPERN*
 R. SCOTT WEIDE
 MICHAEL K. FRIEDLAND
 DALE C. HUNT
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 JENNY G. KO
 KAROLINE A. DELANEY
 JOHN W. HOLCOMB
 JAMES J. MULLEN, III

OF COUNSEL
 JERRY R. SEILER
 JAPANESE PATENT ATTY
 KATSUHIRO ARAI
 EUROPEAN PATENT ATTY
 MARTIN HELLEBRANDT
 KOREAN PATENT ATTY
 MINCHEOL KIM
 CHINESE PATENT ATTY
 JIAWEI HUANG
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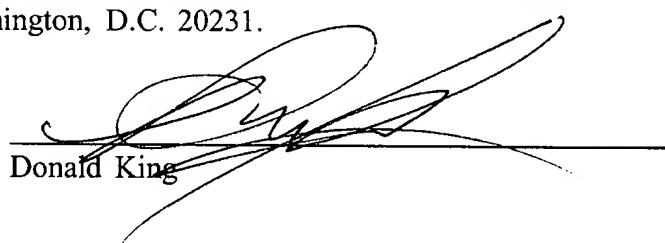


Attorney Docket No. : MASIMO.007C3
Applicant(s) : Diab, Kiani-Azarbajany and Weber
For : SIGNAL PROCESSING APPARATUS
Attorney : Steven C. Jensen
"Express Mail"
Mailing Label No. : EM410970204US
Date of Deposit : July 7, 1998

I hereby certify that the accompanying

Transmittal in Duplicate; Specification in 122 pages and Appendix A in 10 pages; 10 sheets of drawings; **SIGNED** Declaration and Power of Attorney in 3 pages **COPY**; Preliminary Amendment in 2 pages; Request for Interference (5 pages) with attachment - Count in 1 page; Check for Filing Fee(s); Return Prepaid Postcard

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ATTENTION: APPLICATION BRANCH

Sir:

Transmitted herewith for filing is the patent application of

Inventor(s): **Mohamed K. Diab, Esmaiel Kiani-Azarbajany and Walter M. Weber**

For: **SIGNAL PROCESSING APPARATUS**

Enclosed are:

(X) 20 sheets of drawings.

(X) This application is a continuation of prior application no. 08/943,511, filed October 6, 1997, which is a continuation of application no. 08/572,488, filed December 14, 1995 (now U.S. Patent No. 5,685,299), which is a continuation of application no. 08/132,812, filed October 6, 1993 (now U.S. Patent No. 5,490,505).

(X) A copy of Declaration from prior application is enclosed.

(X) Preliminary Amendment is enclosed.

(X) Incorporation by Reference. The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

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FOR	NUMBER FILED	NUMBER EXTRA	RATE	FEE
Basic Fee			\$790	\$790
Total Claims	1 - 20 =	0 x	\$22	\$0
Independent Claims	1 - 3 =	0 x	\$82	\$0
If application contains any multiple dependent claims(s), then add			\$270	\$0
			TOTAL FILING FEE	\$790

(X) Request for Interference in five (5) pages with attachment - Count in one (1) pages.

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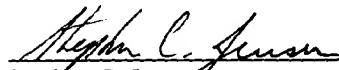
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Stephen C. Jensen
Stephen C. Jensen
Registration No. 35,556
Attorney of Record

SCJ-7461:is/vml
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PATENT

MAS-102 007C3

SIGNAL PROCESSING APPARATUS

Field of the Invention

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The present invention relates to the field of signal processing. More specifically, the present invention relates to the processing of measured signals, containing a primary and a secondary signal, for the removal or derivation of either the primary or secondary signal when little is known about either of these components. The present invention also relates to the use of a novel processor which in conjunction with a correlation canceler, such as an adaptive noise canceler, produces primary and/or secondary signals. The present invention is especially useful for physiological monitoring systems including blood oxygen saturation.

Background of the Invention

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Signal processors are typically employed to remove or derive either the primary or secondary signal portion from a composite measured signal including a primary signal portion and a secondary signal portion. If the secondary signal portion occupies a different frequency spectrum than the primary signal portion, then conventional filtering techniques such as low pass, band pass, and high pass filtering could be used to remove or derive either the primary or the secondary signal portion from the total signal. Fixed single or multiple notch filters could also be employed if the primary and/or secondary signal portion(s) exit at a fixed frequency(s).

It is often the case that an overlap in frequency spectrum between the primary and secondary signal portions exists. Complicating matters further, the statistical properties of one or both of the primary and secondary signal portions change with time. In such cases, conventional filtering techniques are totally ineffective in extracting either the primary or secondary signal. If, however, a description of either the primary or secondary signal portion can be made available correlation canceling, such as adaptive noise canceling, can be employed to remove either the primary or secondary signal portion of the signal leaving the other portion available for measurement.

Correlation cancelers, such as adaptive noise cancelers, dynamically change their transfer function to adapt to and remove either the primary or secondary signal portions of a composite signal. Correlation cancelers require either a secondary reference or a primary reference which is correlated to either the secondary signal or the primary signal portions only. The reference signals are not necessarily a representation of the primary or secondary signal portions, but have a frequency spectrum which is similar to that of the primary or secondary signal portions. In many cases, it requires considerable ingenuity to determine a reference signal since nothing is usually known a priori about the secondary and/or primary signal portions.

One area where composite measured signals comprising a primary signal portion and a secondary signal portion about which no information can easily be determined is physiological monitoring. Physiological monitoring apparatuses generally measure signals derived from a physiological system, such as the human body. Measurements

which are typically taken with physiological monitoring systems include electrocardiographs, blood pressure, blood gas saturation (such as oxygen saturation), capnographs, heart rate, respiration rate, and depth of anesthesia, for example. Other types of measurements include those which measure the pressure and quantity of a substance within the body such as breathalyzer testing, drug testing, cholesterol testing, glucose testing, arterial carbon dioxide testing, protein testing, and carbon monoxide testing, for example. Complications arising in these measurements are often due to motion of the patient, both external and internal (muscle movement, for example), during the measurement process.

Knowledge of physiological systems, such as the amount of oxygen in a patient's blood, can be critical, for example during surgery. These data can be determined by a lengthy invasive procedure of extracting and testing matter, such as blood, from a patient, or by more expedient, non-invasive measures. Many types of non-invasive measurements can be made by using the known properties of energy attenuation as a selected form of energy passes through a medium.

Energy is caused to be incident on a medium either derived from or contained within a patient and the amplitude of transmitted or reflected energy is then measured. The amount of attenuation of the incident energy caused by the medium is strongly dependent on the thickness and composition of the medium through which the energy must pass as well as the specific form of energy selected. Information about a physiological system can be derived from data taken from the attenuated signal of the incident energy transmitted through the medium if either the primary

5 or secondary signal of the composite measurement signal can be removed. However, non-invasive measurements often do not afford the opportunity to selectively observe the interference causing either the primary or secondary signal portions, making it difficult to extract either one of them from the composite signal.

10 The primary and/or secondary signal portions often originate from both AC and/or DC sources. The DC portions are caused by transmission of the energy through differing media which are of relatively constant thickness within the body, such as bone, tissue, skin, blood, etc. These portions are easy to remove from a composite signal. The AC components are caused by physiological pulsations or when differing media being measured are perturbed and thus, change in thickness while the measurement is being made. Since most materials in and derived from the body are easily compressed, the thickness of such matter changes if the patient moves during a non-invasive physiological measurement. Patient movement, muscular movement and vessel movement, can cause the properties of energy attenuation to vary erratically. Traditional signal filtering techniques are frequently totally ineffective and grossly deficient in removing these motion induced effects from a signal. The erratic or unpredictable nature of motion induced signal components is the major obstacle in removing or deriving them. Thus, presently available physiological monitors generally become totally inoperative during time periods when the measurement site is perturbed.

25 30 A blood gas monitor is one example of a physiological monitoring system which is based upon the measurement of energy attenuated by biological tissues or substances. Blood gas monitors transmit light into the tissue and

measure the attenuation of the light as a function of time. The output signal of a blood gas monitor which is sensitive to the arterial blood flow contains a component which is a waveform representative of the patient's arterial pulse. This type of signal, which contains a component related to the patient's pulse, is called a plethysmographic wave, and is shown in Figure 1 as curve s. Plethysmographic waveforms are used in blood pressure or blood gas saturation measurements, for example. As the heart beats, the amount of blood in the arteries increases and decreases, causing increases and decreases in energy attenuation, illustrated by the cyclic wave s in Figure 1.

Typically, a digit such as a finger, an ear lobe, or other portion of the body where blood flows close to the skin, is employed as the medium through which light energy is transmitted for blood gas attenuation measurements. The finger comprises skin, fat, bone, muscle, etc., shown schematically in Figure 2, each of which attenuates energy incident on the finger in a generally predictable and constant manner. However, when fleshy portions of the finger are compressed erratically, for example by motion of the finger, energy attenuation becomes erratic.

An example of a more realistic measured waveform S is shown in Figure 3, illustrating the effect of motion. The primary plethysmographic waveform portion of the signal s is the waveform representative of the pulse, corresponding to the sawtooth-like pattern wave in Figure 1. The large, secondary motion-induced excursions in signal amplitude hide the primary plethysmographic signal s. It is easy to see how even small variations in amplitude make it difficult to distinguish the primary signal s in the presence of a secondary signal component n.

A specific example of a blood gas monitoring apparatus
is a pulse oximeter which measures the arterial saturation
of oxygen in the blood. The pumping of the heart forces
freshly oxygenated blood into the arteries causing greater
energy attenuation. The arterial saturation of oxygenated
blood may be determined from the depth of the valleys
relative to the peaks of two plethysmographic waveforms
measured at separate wavelengths. Patient movement
introduces signal portions mostly due to venous blood, or
5 motion artifacts, to the plethysmographic waveform
illustrated in Figure 3. It is these motion artifacts which
must be removed from the measured signal for the oximeter
to continue the measurement of arterial blood oxygen
saturation, even during periods when the patient moves. It
10 is also these motion artifacts which must be derived from
the measured signal for the oximeter to obtain an estimate
of venous blood oxygen saturation. Once the signal
components due to either arterial blood or venous blood is
known, its corresponding oxygen saturation may be
15 determined.

Summary of the Invention

This invention is an improvement of U. S. Patent
25 Application No. 07/666,060 filed March 7, 1991 and entitled
Signal Processing Apparatus and Method, which earlier
application has been assigned to the assignee of the
instant application. The invention is a signal processor
30 which acquires a first signal and a second signal that is
correlated to the first signal. The first signal comprises
a first primary signal portion and a first secondary signal
portion. The second signal comprises a second primary
signal portion and a second secondary signal portion. The
signals may be acquired by propagating energy through a

medium and measuring an attenuated signal after transmission or reflection. Alternatively, the signals may be acquired by measuring energy generated by the medium.

5 The first and second measured signals are processed to generate a secondary reference which does not contain the primary signal portions from either of the first or second measured signals. The remaining secondary signal portions from the first and second measured signals are combined to form the secondary reference. This secondary reference is correlated to the secondary signal portion of each of the first and second measured signals.

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30 The secondary reference is then used to remove the secondary portion of each of the first and second measured signals via a correlation canceler, such as an adaptive noise canceler. The correlation canceler is a device which takes a first and second input and removes from the first input all signal components which are correlated to the second input. Any unit which performs or nearly performs this function is herein considered to be a correlation canceler. An adaptive correlation canceler can be described by analogy to a dynamic multiple notch filter which dynamically changes its transfer function in response to a reference signal and the measured signals to remove frequencies from the measured signals that are also present in the reference signal. Thus, a typical adaptive correlation canceler receives the signal from which it is desired to remove a component and a reference signal. The output of the correlation canceler is a good approximation to the desired signal with the undesired component removed.

 Alternatively, the first and second measured signals may be processed to generate a primary reference which does

not contain the secondary signal portions from either of the first or second measured signals. The remaining primary signal portions from the first and second measured signals are combined to form the primary reference. The primary reference may then be used to remove the primary portion of each of the first and second measured signals via a correlation canceler. The output of the correlation canceler is a good approximation to the secondary signal with the primary signal removed and may be used for subsequent processing in the same instrument or an auxiliary instrument. In this capacity, the approximation to the secondary signal may be used as a reference signal for input to a second correlation canceler together with either the first or second measured signals for computation of, respectively, either the first or second primary signal portions.

Physiological monitors can often advantageously employ signal processors of the present invention. Often in physiological measurements a first signal comprising a first primary portion and a first secondary portion and a second signal comprising a second primary portion and a second secondary portion are acquired. The signals may be acquired by propagating energy through a patient's body (or a material which is derived from the body, such as breath, blood, or tissue, for example) or inside a vessel and measuring an attenuated signal after transmission or reflection. Alternatively, the signal may be acquired by measuring energy generated by a patient's body, such as in electrocardiography. The signals are processed via the signal processor of the present invention to acquire either a secondary reference or a primary reference which is input to a correlation canceler, such as an adaptive noise canceler.

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One physiological monitoring apparatus which can advantageously incorporate the features of the present invention is a monitoring system which determines a signal which is representative of the arterial pulse, called a plethysmographic wave. This signal can be used in blood pressure calculations, blood gas saturation measurements, etc. A specific example of such a use is in pulse oximetry which determines the saturation of oxygen in the blood. In this configuration, we define the primary portion of the signal to be the arterial blood contribution to attenuation of energy as it passes through a portion of the body where blood flows close to the skin. The pumping of the heart causes blood flow to increase and decrease in the arteries in a periodic fashion, causing periodic attenuation wherein the periodic waveform is the plethysmographic waveform representative of the arterial pulse. We define the secondary portion of the signal to be that which is usually considered to be noise. This portion of the signal is related to the venous blood contribution to attenuation of energy as it passes through the body. Patient movement causes this component to flow in an unpredictable manner, causing unpredictable attenuation and corrupting the otherwise periodic plethysmographic waveform. Respiration also causes secondary or noise component to vary, although typically at a much lower frequency than the patients pulse rate.

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A physiological monitor particularly adapted to pulse oximetry oxygen saturation measurement comprises two light emitting diodes (LED's) which emit light at different wavelengths to produce first and second signals. A detector registers the attenuation of the two different energy signals after each passes through an absorptive media, for example a digit such as a finger, or an earlobe. The

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attenuated signals generally comprise both primary and secondary signal portions. A static filtering system, such as a bandpass filter, removes a portion of the secondary signal which is outside of a known bandwidth of interest, leaving an erratic or random secondary signal portion, often caused by motion and often difficult to remove, along with the primary signal portion.

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DIGITAL DESIGN
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Next, a processor of the present invention removes the primary signal portions from the measured signals yielding a secondary reference which is a combination of the remaining secondary signal portions. The secondary reference is correlated to both of the secondary signal portions. The secondary reference and at least one of the measured signals are input to a correlation canceler, such as an adaptive noise canceler, which removes the random or erratic portion of the secondary signal. This yields a good approximation to the primary plethysmographic signal as measured at one of the measured signal wavelengths. As is known in the art, quantitative measurements of the amount of oxygenated arterial blood in the body can be determined from the plethysmographic signal in a variety of ways.

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The processor of the present invention may also remove the secondary signal portions from the measured signals yielding a primary reference which is a combination of the remaining primary signal portions. The primary reference is correlated to both of the primary signal portions. The primary reference and at least one of the measured signals are input to a correlation canceler which removes the primary portions of the measured signals. This yields a good approximation to the secondary signal at one of the measured signal wavelengths. This signal may be useful for removing secondary signals from an auxiliary instrument as

well as determining venous blood oxygen saturation.

One aspect of the present invention is a signal processor comprising a detector for receiving a first signal which travels along a first propagation path and a second signal which travels along a second propagation path wherein a portion of the first and second propagation paths are located in a propagation medium. The first signal has a first primary signal portion and a first secondary signal portion and the second signal has a second primary signal portion and a second secondary signal portion. The first and second secondary signal portions are a result of a change of the propagation medium. This aspect of the invention additionally comprises a reference processor having an input for receiving the first and second signals. The processor is adapted to combine the first and second signals to generate a secondary reference having a significant component which is a function of the first and said second secondary signal portions. The processor may also be adapted to combine the first and second signals to generate a primary reference having a significant component which is a function of the first and second primary signal portions

The above described aspect of the present invention may further comprise a signal processor for receiving the secondary reference signal and the first signal and for deriving therefrom an output signal having a significant component which is a function of the first primary signal portion of the first signal. Alternatively, the above described aspect of the present invention may further comprise a signal processor for receiving the secondary reference signal and the second signal and for deriving therefrom an output signal having a significant component

which is a function of the second primary signal portion of
the second signal. Alternatively, the above described
aspect of the present invention may further comprise a
signal processor for receiving the primary reference and
the first signal and for deriving therefrom an output
signal having a significant component which is a function
of the first secondary signal portion of the signal of the
first signal. Alternatively, the above described aspect of
the present invention may further comprise a signal
processor for receiving the primary reference and the
second signal and for deriving therefrom an output signal
having a significant component which is a function of the
second secondary signal portion of the second signal. The
signal processor may comprise a correlation canceler, such
as an adaptive noise canceler. The adaptive noise canceler
may comprise a joint process estimator having a least-
squares-lattice predictor and a regression filter.

The detector in the aspect of the signal processor of
the present invention described above may further comprise
a sensor for sensing a physiological function. The sensor
may comprise a light or other electromagnetic sensitive
device. Additionally, the present invention may further
comprise a pulse oximeter for measuring oxygen saturation
in a living organism. The present invention may further
comprise an electrocardiograph.

Another aspect of the present invention is a
physiological monitoring apparatus comprising a detector
for receiving a first physiological measurement signal
which travels along a first propagation path and a second
physiological measurement signal which travels along a
second propagation path. A portion of the first and second
propagation paths being located in the same propagation

medium. The first signal has a first primary signal portion and a first secondary signal portion and the second signal has a second primary signal portion and a second secondary signal portion. The physiological monitoring apparatus further comprises a reference processor having an input for receiving the first and second signals. The processor is adapted to combine the first and second signals to generate a secondary reference signal having a significant component which is a function of the first and the second secondary signal portions. Alternatively, the processor may be adapted to combine the first and second signals to generate a primary reference having a component which is a function of the first and second primary signal portions.

The physiological monitoring apparatus may further comprise a signal processor for receiving the secondary reference and the first signal and for deriving therefrom an output signal having a significant component which is a function of the first primary signal portion of the first signal. Alternatively, the physiological monitoring apparatus may further comprise a signal processor for receiving the secondary reference and the second signal and for deriving therefrom an output signal having a significant component which is a function of the second primary signal portion of the second signal. Alternatively, the physiological monitoring apparatus may further comprise a signal processor for receiving the primary reference and the first signal and deriving therefrom an output signal having a significant component which is a function of the first secondary signal portion of the first signal. Alternatively, the physiological monitoring apparatus may further comprise a signal processor for receiving the primary reference and the second signal and deriving therefrom an output signal having a significant component

which is a function of the second secondary signal portion of the second signal.

A further aspect of the present invention is an apparatus for measuring a blood constituent comprising an energy source for directing a plurality of predetermined wavelengths of electromagnetic energy upon a specimen and a detector for receiving the plurality of predetermined wavelengths of electromagnetic energy from the specimen. The detector produces electrical signals corresponding to the predetermined wavelengths in response to the electromagnetic energy. At least two of the electrical signals are used each having a primary signal portion and an secondary signal portion. Additionally, the apparatus comprises a reference processor having an input for receiving the electrical signals. The processor is configured to combine said electrical signals to generate a secondary reference having a significant component which is derived from the secondary signal portions. Alternatively, the processor may be configured to combine said signals to generate a primary reference having a significant component which is derived from the primary signal portions.

This aspect of the present invention may further comprise a signal processor for receiving the secondary reference and one of the two electrical signals and for deriving therefrom an output signal having a significant component which is a function of the primary signal portion of one of the two electrical signals. Another aspect of the present invention may further comprise a signal processor for receiving the primary reference and one of the two electrical signals and for deriving therefrom an output signal having a significant component which is a function of the secondary signal portion of one of the two

5 electrical signals. This may be accomplished by use of a correlation canceler, such as an adaptive noise canceler, in the signal processor which may employ a joint process estimator having a least-squares-lattice predictor and a regression filter.

10 Yet another aspect of the present invention is a blood gas monitor for non-invasively measuring a blood constituent in a body comprising a light source for directing at least two predetermined wavelengths of light upon a body and a detector for receiving the light from the body. The detector, in response to the light from the body, produces at least two electrical signals corresponding to the at least two predetermined wavelengths of light. The at least two electrical signals each have a primary signal portion and a secondary signal portion. The blood oximeter further comprises a reference processor having an input for receiving the at least two electrical signals. The processor is adapted to combine the at least two electrical signals to generate a secondary reference with a significant component which is derived from the secondary signal portions. The blood oximeter may further comprise a signal processor for receiving the secondary reference and the two electrical signals and for deriving therefrom at least two output signals which are substantially equal, respectively, to the primary signal portions of the electrical signals. Alternatively, the reference processor may be adapted to combine the at least two electrical signals to generate a primary reference with a significant component which is derived from the primary signal portions. The blood oximeter may further comprise a signal processor for receiving the primary reference and the two electrical signals and for deriving therefrom at least two output signals which are substantially equivalent to the

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secondary signal portions of the electrical signal. The signal processor may comprise a joint process estimator.

The present invention also includes a method of determining a secondary reference from a first signal comprising a first primary signal portion and a first secondary portion and a second signal comprising a second primary signal portion and a second secondary portion. The method comprises the steps of selecting a signal coefficient which is proportional to a ratio of predetermined attributes of the first primary signal portion and predetermined attributes of the second primary signal portion. The first signal and the signal coefficient are input into a signal multiplier wherein the first signal is multiplied by the signal coefficient thereby generating a first intermediate signal. The second signal and the first intermediate signal are input into a signal subtractor wherein the first intermediate signal is subtracted from the second signal. This generates a secondary reference having a significant component which is derived from the first and second secondary signal portions.

The present invention also includes a method of determining a primary reference from a first signal comprising a first primary signal portion and a first secondary signal portion and a second signal comprising a second primary signal portion and a second secondary signal portion. The method comprises the steps of selecting a signal coefficient which is proportional to a ratio of the predetermined attributes of the first secondary signal portion and predetermined attributes of the second secondary signal portion. The first signal and the signal coefficient are input into a signal multiplier wherein the

first signal is multiplied by the signal coefficient thereby generating a first intermediate signal. The second signal and the first intermediate signal are input into a signal subtractor wherein the first intermediate signal is subtracted from the second signal. This generates a primary reference having a significant component which is derived from the first and second primary signal portions. The first and second signals in this method may be derived from electromagnetic energy transmitted through an absorbing medium.

The present invention further embodies a physiological monitoring apparatus comprising means for acquiring a first signal comprising a first primary signal portion and a first secondary signal portion and a second signal comprising a second primary signal portion and a second secondary signal portion. The physiological monitoring apparatus of the present invention also comprises means for determining from the first and second signals a secondary reference. Additionally, the monitoring apparatus comprises a correlation canceler, such as an adaptive noise canceler, having a secondary reference input for receiving the secondary reference and a signal input for receiving the first signal wherein the correlation canceler, in real or near real time, generates an output signal which approximates the first primary signal portion. Alternatively, the physiological monitoring device may also comprise means for determining from the first and second signals a primary reference. Additionally, the monitoring apparatus comprises a correlation canceler having a primary reference input for receiving the primary reference and a signal input for receiving the first signal wherein the correlation canceler, in real or near real time, generates an output signal which approximates the first secondary

signal portion. The correlation canceler may further comprise a joint process estimator.

A further aspect of the present invention is an apparatus for processing an amplitude modulated signal having a signal amplitude complicating feature, the apparatus comprising an energy source for directing electromagnetic energy upon a specimen. Additionally, the apparatus comprises a detector for acquiring a first amplitude modulated signal and a second amplitude modulated signal. Each of the first and second signals has a component containing information about the attenuation of electromagnetic energy by the specimen and a signal amplitude complicating feature. The apparatus includes a reference processor for receiving the first and second amplitude modulated signals and deriving therefrom a secondary reference which is correlated with the signal amplitude complicating feature. Further, the apparatus incorporates a correlation canceler having a signal input for receiving the first amplitude modulated signal, a secondary reference input for receiving the secondary reference, wherein the correlation canceler produces an output signal having a significant component which is derived from the component containing information about the attenuation of electromagnetic energy by the specimen. Alternatively, the apparatus may also include a reference processor for receiving the first and second amplitude modulated signals and deriving therefrom a primary reference which is correlated with the component containing information about the attenuation of electromagnetic energy by the specimen. Further, the apparatus incorporates a correlation canceler having a signal input for receiving the first amplitude modulated signal, a primary reference input for receiving the primary reference, wherein the

correlation canceler produces an output signal having a primary component which is derived from the signal amplitude complicating feature.

Still another aspect of the present invention is an apparatus for extracting a plethysmographic waveform from an amplitude modulated signal having a signal amplitude complicating feature, the apparatus comprising a light source for transmitting light into an organism and a detector for monitoring light from the organism. The detector produces a first light attenuation signal and a second light attenuation signal, wherein each of the first and second light attenuation signals has a component which is representative of a plethysmographic waveform and a component which is representative of the signal amplitude complicating feature. The apparatus also includes a reference processor for receiving the first and second light attenuation signals and deriving therefrom a secondary reference. The secondary reference and the signal amplitude complicating feature each have a frequency spectrum. The frequency spectrum of the secondary reference is correlated with the frequency spectrum of the signal amplitude complicating feature. Additionally incorporated into this embodiment of the present invention is a correlation canceler having a signal input for receiving the first attenuation signal and a secondary reference input for receiving the secondary reference. The correlation canceler produces an output signal having a significant component which is derived from the component which is representative of a plethysmographic waveform. The apparatus may also include a reference processor for receiving the first and second light attenuation signals and deriving therefrom a primary reference. Additionally incorporated in this embodiment of the present invention is

5 a correlation canceler having a signal input for receiving
the first attenuation signal and a primary reference input
for receiving the primary reference. The correlation
canceler produces an output signal having a significant
component which is derived from the component which is
representative of the signal complicating feature.

10 The present invention also comprises a method of
removing or determining a motion artifact signal from a
signal derived from a physiological measurement wherein a
first signal having a physiological measurement component
and a motion artifact component and a second signal having
a physiological measurement component and a motion artifact
component are acquired. From the first and second signals a
secondary reference which is a primary function of the
first and second signals motion artifact components is
derived. This method of removing a motion artifact signal
from a signal derived from a physiological measurement may
also comprise the step of inputting the secondary reference
into a correlation canceler, such as an adaptive noise
canceler, to produce an output signal which is a
significant function of the physiological measurement
component of the first or second signal. Alternatively,
from the first and second signals a primary reference which
is a significant function of the physiological measurement
components of the first and second signals may be derived.
This approach may also comprise the step of inputting the
primary reference into a correlation canceler to produce an
output signal which is a significant function of the first
or second signal's motion artifact component.

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Brief Description of the Drawings

Figure 1 illustrates an ideal plethysmographic waveform.

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Figure 2 schematically illustrates the cross-sectional structure of a typical finger.

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Figure 3 illustrates a plethysmographic waveform which includes a motion-induced erratic signal portion.

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Figure 4a illustrates a schematic diagram of a physiological monitor, to compute primary physiological signals, incorporating a processor of the present invention, and a correlation canceler.

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Figure 4b illustrates a schematic diagram of a physiological monitor, to compute secondary erratic signals, incorporating a processor of the present invention, and a correlation canceler.

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Figure 5a illustrates an example of an adaptive noise canceler which could be employed in a physiological monitor, to compute primary physiological signals, which also incorporates the processor of the present invention.

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Figure 5b illustrates an example of an adaptive noise canceler which could be employed in a physiological monitor, to compute secondary motion artifact signals, which also incorporates the processor of the present invention.

Figure 5c illustrates the transfer function of a multiple notch filter.

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Figure 6a illustrates a schematic absorbing material comprising N constituents within an absorbing material.

Figure 6b illustrates another schematic absorbing material comprising N constituents, including one mixed layer, within an absorbing material.

Figure 6c illustrates another schematic absorbing material comprising N constituents, including two mixed layers, within an absorbing material.

Figure 7a illustrates a schematic diagram of a monitor, to compute primary and secondary signals, incorporating a processor of the present invention, a plurality of signal coefficients $\omega_1, \omega_2, \dots, \omega_n$, and a correlation canceler.

Figure 7b illustrates the ideal correlation canceler energy or power output as a function of the signal coefficients $\omega_1, \omega_2, \dots, \omega_n$. In this particular example, $\omega_3 = \omega_a$ and $\omega_7 = \omega_v$.

Figure 7c illustrates the non-ideal correlation canceler energy or power output as a function of the signal coefficients $\omega_1, \omega_2, \dots, \omega_n$. In this particular example, $\omega_3 = \omega_a$ and $\omega_7 = \omega_v$.

Figure 8 is a schematic model of a joint process estimator comprising a least-squares lattice predictor and a regression filter.

Figure 9 is a flowchart representing a subroutine capable of implementing a joint process estimator as modeled in Figure 8.

Figure 10 is a schematic model of a joint process estimator with a least-squares lattice predictor and two regression filters.

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Figure 11 is an example of a physiological monitor incorporating a processor of the present invention and a correlation canceler within a microprocessor. This physiological monitor is specifically designed to measure a plethysmographic waveform or a motion artifact waveform and perform oximetry measurements.

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Figure 12 is a graph of oxygenated and deoxygenated hemoglobin absorption coefficients vs. wavelength.

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Figure 13 is a graph of the ratio of the absorption coefficients of deoxygenated hemoglobin divided by oxygenated hemoglobin vs. wavelength.

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Figure 14 is an expanded view of a portion of Figure 12 marked by a circle labeled 13.

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Figure 15 illustrates a signal measured at a first red wavelength $\lambda_a = \lambda_{red1} = 650\text{nm}$ for use in a processor of the present invention employing the ratiometric method for determining either the primary reference $n'(t)$ or the secondary reference $s'(t)$ and for use in a correlation canceler, such as an adaptive noise canceler. The measured signal comprises a primary portion $s_{\lambda_a}(t)$ and a secondary portion $n_{\lambda_a}(t)$.

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Figure 16 illustrates a signal measured at a second red wavelength $\lambda_b = \lambda_{red2} = 685\text{nm}$ for use in a processor of the present invention employing the ratiometric method for

5 determining the secondary reference $n'(t)$ or the primary reference $s'(t)$. The measured signal comprises a primary portion $s_{\lambda b}(t)$ and a secondary portion $n_{\lambda b}(t)$.

10 Figure 17 illustrates a signal measured at an infrared wavelength $\lambda_c = \lambda_{IR} = 940\text{nm}$ for use in a correlation canceler. The measured signal comprises a primary portion $s_{\lambda c}(t)$ and a secondary portion $n_{\lambda c}(t)$.

15 Figure 18 illustrates the secondary reference $n'(t)$ determined by a processor of the present invention using the ratiometric method.

20 Figure 19 illustrates the primary reference $s'(t)$ determined by a processor of the present invention using the ratiometric method.

25 Figure 20 illustrates a good approximation $s''_{\lambda a}(t)$ to the primary portion $s_{\lambda a}(t)$ of the signal $S_{\lambda a}(t)$ measured at $\lambda_a = \lambda_{red1} = 650\text{nm}$ estimated by correlation cancellation with a secondary reference $n'(t)$ determined by the ratiometric method.

30 Figure 21 illustrates a good approximation $s''_{\lambda c}(t)$ to the primary portion $s_{\lambda c}(t)$ of the signal $S_{\lambda c}(t)$ measured at $\lambda_c = \lambda_{IR} = 940\text{nm}$ estimated by correlation cancellation with a secondary reference $n'(t)$ determined by the ratiometric method.

35 Figure 22 illustrates a good approximation $n''_{\lambda a}(t)$ to the secondary portion $n_{\lambda a}(t)$ of the signal $S_{\lambda a}(t)$ measured at $\lambda_a = \lambda_{red1} = 650\text{nm}$ estimated by correlation cancellation with a primary reference $s'(t)$ determined by the ratiometric method.

Figure 23 illustrates a good approximation $n''\lambda_c(t)$ to the secondary portion $n\lambda_c(t)$ of the signal $S\lambda_c(t)$ measured at $\lambda_c=\lambda_{IR}=940\text{nm}$ estimated by correlation cancelation with a primary reference $s'(t)$ determined by the ratiometric method.

Figure 24 illustrates a signal measured at a red wavelength $\lambda_a=\lambda_{red}=660\text{nm}$ for use in a processor of the present invention employing the constant saturation method for determining the secondary reference $n'(t)$ or the primary reference $s'(t)$ and for use in a correlation canceler. The measured signal comprises a primary portion $S\lambda_a(t)$ and a secondary portion $n\lambda_a(t)$.

Figure 25 illustrates a signal measured at an infrared wavelength $\lambda_b=\lambda_{IR}=940\text{nm}$ for use in a processor of the present invention employing the constant saturation method for determining the secondary reference $n'(t)$ or the primary reference $s'(t)$ and for use in a correlation canceler. The measured signal comprises a primary portion $S\lambda_b(t)$ and a secondary portion $n\lambda_b(t)$.

Figure 26 illustrates the secondary reference $n'(t)$ determined by a processor of the present invention using the constant saturation method.

Figure 27 illustrates the primary reference $s'(t)$ determined by a processor of the present invention using the constant saturation method.

Figure 28 illustrates a good approximation $s''\lambda_a(t)$ to the primary portion $S\lambda_a(t)$ of the signal $S\lambda_a(t)$ measured at $\lambda_a=\lambda_{red}=660\text{nm}$ estimated by correlation cancelation with a

secondary reference $n'(t)$ determined by the constant saturation method.

Figure 29 illustrates a good approximation $s''\lambda_b(t)$ to the primary portion $s_{\lambda_b}(t)$ of the signal $S_{\lambda_b}(t)$ measured at $\lambda_b=\lambda_{IR}=940\text{nm}$ estimated by correlation cancelation with a secondary reference $n'(t)$ determined by the constant saturation method.

Figure 30 illustrates a good approximation $n''\lambda_a(t)$ to the secondary portion $n_{\lambda_a}(t)$ of the signal $S_{\lambda_a}(t)$ measured at $\lambda_a=\lambda_{red}=660\text{nm}$ estimated by correlation cancelation with a primary reference $s'(t)$ determined by the constant saturation method.

Figure 31 illustrates a good approximation $n''\lambda_b(t)$ to the secondary portion $n_{\lambda_b}(t)$ of the signal $S_{\lambda_b}(t)$ measured at $\lambda_b=\lambda_{IR}=940\text{nm}$ estimated by correlation cancelation with a primary reference $s'(t)$ determined by the constant saturation method.

Figure 32 depicts a set of 3 concentric electrodes, i. e. a tripolar electrode sensor, to derive electrocardiography (ECG) signals, denoted as S_1 , S_2 and S_3 , for use with the present invention. Each of the ECG signals contains a primary portion and a secondary portion.

Detailed Description of the Invention

The present invention is a processor which determines either a secondary reference $n'(t)$ or a primary reference $s'(t)$ for use in a correlation canceler, such as an adaptive noise canceler. A correlation canceler may estimate a good approximation $s''(t)$ to a primary signal $s(t)$ from a composite signal $S(t) = s(t) + n(t)$ which, in addition to the primary portion $s(t)$ comprises a secondary portion $n(t)$. It may also be used to provide a good approximation $n''(t)$ to the secondary signal $n(t)$. The secondary portion $n(t)$ may contain one or more of a constant portion, a predictable portion, an erratic portion, a random portion, etc. The approximation to the primary signal $s''(t)$ or secondary signal $n''(t)$ is derived by removing as many of the secondary portions $n(t)$ or primary portions $s(t)$ from the composite signal $S(t)$ as possible. The constant portion and predictable portion are easily removed with traditional filtering techniques, such as simple subtraction, low pass, band pass, and high pass filtering. The erratic portion is more difficult to remove due to its unpredictable nature. If something is known about the erratic signal, even statistically, it could be removed, at least partially, from the measured signal via traditional filtering techniques. However, it is often the case that no information is known about the erratic portion of the noise. In this case, traditional filtering techniques are usually insufficient. Often no information about the erratic portion of the measured signal is known. Thus, a correlation canceler, such as an adaptive noise canceler may be utilized in the present invention to remove or derive the erratic portion.

Generally, a correlation canceler has two signal

inputs and one output. One of the inputs is either the secondary reference $n'(t)$ or the primary reference $s'(t)$ which are correlated, respectively, to the secondary signal portions $n(t)$ and the primary signal portions $s(t)$ present in the composite signal $S(t)$. The other input is for the composite signal $S(t)$. Ideally, the output of the correlation canceler $s''(t)$ or $n''(t)$ corresponds, respectively, to the primary signal $s(t)$ or the secondary signal $n(t)$ portions only. Often, the most difficult task in the application of correlation cancelers is determining the reference signals $n'(t)$ and $s'(t)$ which are correlated to the secondary $n(t)$ and primary $s(t)$ portions, respectively, of the measured signal $S(t)$ since, as discussed above, these portions are quite difficult to isolate from the measured signal $S(t)$. In the signal processor of the present invention, either a secondary reference $n'(t)$ or a primary reference $s'(t)$ is determined from two composite signals measured simultaneously, or nearly simultaneously, at two different wavelengths, λ_a and λ_b .

A block diagram of a generic monitor incorporating a signal processor, or reference processor, according to the present invention, and a correlation canceler is shown in Figures 4a and 4b. Two measured signals, $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$, are acquired by a detector 20. One skilled in the art will realize that for some physiological measurements, more than one detector may be advantageous. Each signal is conditioned by a signal conditioner 22a and 22b. Conditioning includes, but is not limited to, such procedures as filtering the signals to remove constant portions and amplifying the signals for ease of manipulation. The signals are then converted to digital data by an analog-to-digital converter 24a and 24b. The

first measured signal $s_{\lambda a}(t)$ comprises a first primary signal portion, labeled herein $s_{\lambda a}(t)$, and a first secondary signal portion, labeled herein $n_{\lambda a}(t)$. The second measured signal $s_{\lambda b}(t)$ is at least partially correlated to the first measured signal $s_{\lambda a}(t)$ and comprises a second primary signal portion, labeled herein $s_{\lambda b}(t)$, and a second secondary signal portion, labeled herein $n_{\lambda b}(t)$. Typically the first and second secondary signal portions, $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$, are uncorrelated and/or erratic with respect to the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$. The secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ are often caused by motion of a patient. The signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ are input to a reference processor 26. The reference processor 26 multiplies the second measured signal $s_{\lambda b}(t)$ by either a factor $\omega_a = s_{\lambda a}(t)/s_{\lambda b}(t)$ or a factor $\omega_v = n_{\lambda a}(t)/n_{\lambda b}(t)$ and then subtracts the second measured signal $s_{\lambda b}(t)$ from the first measured signal $s_{\lambda a}(t)$. The signal coefficient factors ω_a and ω_v are determined to cause either the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ or the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ to cancel when the two signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ are subtracted. Thus, the output of the reference processor 26 is either a secondary reference signal $n'(t) = n_{\lambda a}(t) - \omega_a n_{\lambda b}(t)$, in Figure 4a, which is correlated to both of the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ or a primary reference signal $s'(t) = s_{\lambda a}(t) - \omega_v s_{\lambda b}(t)$, in Figure 4b, which is correlated to both of the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$. A reference signal $n'(t)$ or $s'(t)$ is input, along with one of the measured signals $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$, to a correlation canceler 27 which uses the reference signal $n'(t)$ or $s'(t)$ to remove either the secondary signal portions $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$ or the primary signal portions $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ from the measured signal $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$. The output of the correlation canceler 27 is a good

approximation $s''(t)$ or $n''(t)$ to either the primary $s(t)$ or the secondary $n(t)$ signal components. The approximation $s''(t)$ or $n''(t)$ is displayed on the display 28.

5 An adaptive noise canceler 30, an example of which is shown in block diagram form in Figure 5a, is employed to remove either one of the erratic, secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ from the first and second signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$. The adaptive noise canceler 30, which performs
10 the functions of a correlation canceler, in Figure 5a has as one input a sample of the secondary reference $n'(t)$ which is correlated to the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$. The secondary reference $n'(t)$ is determined from the two measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ by the processor 26 of the present invention as described herein.
15 A second input to the adaptive noise canceler, is a sample of either the first or second composite measured signals $S_{\lambda a}(t) = s_{\lambda a}(t) + n_{\lambda a}(t)$ or $S_{\lambda b}(t) = s_{\lambda b}(t) + n_{\lambda b}(t)$.

20 The adaptive noise canceler 30, in Figure 5b, may also be employed to remove either one of primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ from the first and second signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$. The adaptive noise canceler 30 has as one input a sample of the primary reference $s'(t)$ which is correlated to the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$. The primary reference $s'(t)$ is determined from the two measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ by the processor 26 of the present invention as described herein. A second input to the adaptive noise canceler 30 is a sample of either the first or second measured signals $S_{\lambda a}(t) = s_{\lambda a}(t) + n_{\lambda a}(t)$ or $S_{\lambda b}(t) = s_{\lambda b}(t) + n_{\lambda b}(t)$.
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The adaptive noise canceler 30 functions to remove frequencies common to both the reference $n'(t)$ or $s'(t)$ and

the measured signal $S_{\lambda a}(t)$ or $S_{\lambda b}(t)$. Since the reference signals are correlated to either the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ or the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$, the reference signals will be correspondingly erratic or well behaved. The adaptive noise canceler 30 acts in a manner which may be analogized to a dynamic multiple notch filter based on the spectral distribution of the reference signal $n'(t)$ or $s'(t)$.

Referring to Figure 5c, the transfer function of a multiple notch filter is shown. The notches, or dips in the amplitude of the transfer function, indicate frequencies which are attenuated or removed when a composite measured signal passes through the notch filter. The output of the notch filter is the composite signal having frequencies at which a notch was present removed. In the analogy to an adaptive noise canceler 30, the frequencies at which notches are present change continuously based upon the inputs to the adaptive noise canceler 30.

The adaptive noise canceler 30 shown in Figures 5a and 5b produces an output signal, labeled herein as $s''_{\lambda a}(t)$, $s''_{\lambda b}(t)$, $n''_{\lambda a}(t)$ or $n''_{\lambda b}(t)$ which is fed back to an internal processor 32 within the adaptive noise canceler 30. The internal processor 32 automatically adjusts its own transfer function according to a predetermined algorithm such that the output of the internal processor 32, labeled $b(t)$ in Figure 5a or $c(t)$ in Figure 5b, closely resembles either the secondary signal portion $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$ or the primary signal portion $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$. The output $b(t)$ of the internal processor 32 in Figure 5a is subtracted from the measured signal, $S_{\lambda a}(t)$ or $S_{\lambda b}(t)$, yielding a signal output $s''_{\lambda a}(t) = s_{\lambda a}(t) + n_{\lambda a}(t) - b_{\lambda a}(t)$ or a signal output $s''_{\lambda b}(t) = s_{\lambda b}(t) + n_{\lambda b}(t) - b_{\lambda b}(t)$. The internal

processor optimizes $s''\lambda_a(t)$ or $s''\lambda_b(t)$ such that $s''\lambda_a(t)$ or $s''\lambda_b(t)$ is approximately equal to the primary signal $s_{\lambda_a}(t)$ or $s_{\lambda_b}(t)$, respectively. The output $c(t)$ of the internal processor 32 in Figure 5b is subtracted from the measured signal, $s_{\lambda_a}(t)$ or $s_{\lambda_b}(t)$, yielding a signal output given by
5 $n''\lambda_a(t) = s_{\lambda_a}(t) + n_{\lambda_a}(t) - c_{\lambda_a}(t)$ or a signal output given by $n''\lambda_b(t) = s_{\lambda_b}(t) + n_{\lambda_b}(t) - c_{\lambda_b}(t)$. The internal processor optimizes $n''\lambda_a(t)$ or $n''\lambda_b(t)$ such that $n''\lambda_a(t)$ or
10 $n''\lambda_b(t)$ is approximately equal to the secondary signal $n_{\lambda_a}(t)$ or $n_{\lambda_b}(t)$, respectively.

One algorithm which may be used for the adjustment of the transfer function of the internal processor 32 is a least-squares algorithm, as described in Chapter 6 and Chapter 12 of the book Adaptive Signal Processing by Bernard Widrow and Samuel Stearns, published by Prentice Hall, copyright 1985. This entire book, including Chapters 6 and 12, is hereby incorporated herein by reference.
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Adaptive processors 30 in Figures 5a and 5b have been successfully applied to a number of problems including antenna sidelobe canceling, pattern recognition, the elimination of periodic interference in general, and the elimination of echoes on long distance telephone transmission lines. However, considerable ingenuity is often required to find a suitable reference signal $n'(t)$ or
25 $s'(t)$ since the portions $n_{\lambda_a}(t)$, $n_{\lambda_b}(t)$, $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ cannot easily be separated from the measured signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$. If either the actual secondary portion $n_{\lambda_a}(t)$ or $n_{\lambda_b}(t)$ or the primary signal portion $s_{\lambda_a}(t)$ or $s_{\lambda_b}(t)$
30 were a priori available, techniques such as correlation cancellation would not be necessary. The determination of a suitable reference signal $n'(t)$ or $s'(t)$ from measurements taken by a monitor incorporating a reference processor of

the present invention is one aspect of the present invention.

GENERALIZED DETERMINATION OF PRIMARY AND SECONDARY

5

REFERENCE SIGNALS

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An explanation which describes how the reference signals $n'(t)$ and $s'(t)$ may be determined follows. A first signal is measured at, for example, a wavelength λ_a , by a detector yielding a signal $s_{\lambda_a}(t)$:

$$s_{\lambda_a}(t) = s_{\lambda_a}(t) + n_{\lambda_a}(t) \quad (1)$$

15

where $s_{\lambda_a}(t)$ is the primary signal and $n_{\lambda_a}(t)$ is the secondary signal.

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A similar measurement is taken simultaneously, or nearly simultaneously, at a different wavelength, λ_b , yielding:

$$s_{\lambda_b}(t) = s_{\lambda_b}(t) + n_{\lambda_b}(t). \quad (2)$$

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Note that as long as the measurements, $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$, are taken substantially simultaneously, the secondary signal components, $n_{\lambda_a}(t)$ and $n_{\lambda_b}(t)$, will be correlated because any random or erratic functions will affect each measurement in nearly the same fashion. The well behaved primary signal components, $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$, will also be correlated to one another.

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To obtain the reference signals $n'(t)$ and $s'(t)$, the measured signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ are transformed to eliminate, respectively, the primary or secondary signal

components. One way of doing this is to find proportionality constants, ω_a and ω_v , between the primary signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ and secondary signals $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ such that:

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$$s_{\lambda a}(t) = \omega_a s_{\lambda b}(t)$$

(3)

$$n_{\lambda a}(t) = \omega_v n_{\lambda b}(t).$$

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These proportionality relationships can be satisfied in many measurements, including but not limited to absorption measurements and physiological measurements. Additionally, in most measurements, the proportionality constants ω_a and ω_v can be determined such that:

$$n_{\lambda a}(t) \neq \omega_a n_{\lambda b}(t)$$

(4)

$$s_{\lambda a}(t) \neq \omega_v s_{\lambda b}(t).$$

Multiplying equation (2) by ω_a and then subtracting equation (2) from equation (1) results in a single equation wherein the primary signal terms $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ cancel, leaving:

$$n'(t) = s_{\lambda a}(t) - \omega_a s_{\lambda b}(t) = n_{\lambda a}(t) - \omega_a n_{\lambda b}(t); (5a)$$

a non-zero signal which is correlated to each secondary signal portion $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ and can be used as the secondary reference $n'(t)$ in a correlation canceler such as an adaptive noise canceler.

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Multiplying equation (2) by ω_v and then subtracting equation (2) from equation (1) results in a single equation wherein the secondary signal terms $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$

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cancel, leaving:

$$s'(t) = s_{\lambda a}(t) - \omega_v s_{\lambda b}(t) = s_{\lambda a}(t) - \omega_v s_{\lambda b}(t); \quad (5b)$$

5 a non-zero signal which is correlated to each of the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ and can be used as the signal reference $s'(t)$ in a correlation canceler such as an adaptive noise canceler.

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EXAMPLE OF DETERMINATION OF PRIMARY AND SECONDARY

REFERENCE SIGNALS IN AN ABSORPTIVE SYSTEM

15 Correlation canceling is particularly useful in a large number of measurements generally described as absorption measurements. An example of an absorption type monitor which can advantageously employ correlation canceling, such as adaptive noise canceling, based upon a reference $n'(t)$ or $s'(t)$ determined by a processor of the present invention is one which determines the concentration of an energy absorbing constituent within an absorbing material when the material is subject to change. Such changes can be caused by forces about which information is desired or primary, or alternatively, by random or erratic secondary forces such as a mechanical force on the material. Random or erratic interference, such as motion, generates secondary components in the measured signal. These secondary components can be removed or derived by the correlation canceler if a suitable secondary reference 20 $n'(t)$ or primary reference $s'(t)$ is known.

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35 A schematic N constituent absorbing material comprising a container 42 having N different absorbing constituents, labeled $A_1, A_2, A_3, \dots, A_N$, is shown schematically in Figure 6a. The constituents A_1 through A_N

in Figure 6a are arranged in a generally orderly, layered fashion within the container 42. An example of a particular type of absorptive system is one in which light energy passes through the container 42 and is absorbed according to the generalized Beer-Lambert Law of light absorption. For light of wavelength λ_a , this attenuation may be approximated by:

$$I = I_0 \exp(-\sum_{i=1}^N \varepsilon_i, \lambda_a c_i x_i) \quad (6)$$

Initially transforming the signal by taking the natural logarithm of both sides and manipulating terms, the signal is transformed such that the signal components are combined by addition rather than multiplication, i.e.:

$$S_{\lambda_a} = \ln(I_0/I) = \sum_{i=1}^N \varepsilon_i, \lambda_a c_i x_i \quad (7)$$

where I_0 is the incident light energy intensity; I is the transmitted light energy intensity; ε_i, λ_a is the absorption coefficient of the i^{th} constituent at the wavelength λ_a ; $x_i(t)$ is the optical path length of i^{th} layer, i.e., the thickness of material of the i^{th} layer through which optical energy passes; and $c_i(t)$ is the concentration of the i^{th} constituent in the volume associated with the thickness $x_i(t)$. The absorption coefficients ε_1 through ε_N are known values which are constant at each wavelength. Most concentrations $c_1(t)$ through $c_N(t)$ are typically unknown, as are most of the optical path lengths $x_i(t)$ of each layer. The total optical path length is the sum of each of the individual optical path lengths $x_i(t)$ of each layer.

When the material is not subject to any forces which cause change in the thicknesses of the layers, the optical

path length of each layer, $x_i(t)$, is generally constant. This results in generally constant attenuation of the optical energy and thus, a generally constant offset in the measured signal. Typically, this portion of the signal is of little interest since knowledge about a force which perturbs the material is usually desired. Any signal portion outside of a known bandwidth of interest, including the constant undesired signal portion resulting from the generally constant absorption of the constituents when not subject to change, should be removed. This is easily accomplished by traditional band pass filtering techniques. However, when the material is subject to forces, each layer of constituents may be affected by the perturbation differently than each other layer. Some perturbations of the optical path lengths of each layer $x_i(t)$ may result in excursions in the measured signal which represent desired or primary information. Other perturbations of the optical path length of each layer $x_i(t)$ cause undesired or secondary excursions which mask primary information in the measured signal. Secondary signal components associated with secondary excursions must also be removed to obtain primary information from the measured signal. Similarly, the ability to compute secondary signal components caused by secondary excursions directly allows one to obtain primary signal components from the measured signal via simple subtraction, or correlation cancellation techniques.

The correlation canceler may selectively remove from the composite signal, measured after being transmitted through or reflected from the absorbing material, either the secondary or the primary signal components caused by forces which perturb or change the material differently from the forces which perturbed or changed the material to cause respectively, either the primary or secondary signal

component. For the purposes of illustration, it will be assumed that the portion of the measured signal which is deemed to be the primary signal $s_{\lambda a}(t)$ is the attenuation term $\epsilon_5 c_5 x_5(t)$ associated with a constituent of interest, namely A_5 , and that the layer of constituent A_5 is affected by perturbations different than each of the layers of other constituents A_1 through A_4 and A_6 through A_N . An example of such a situation is when layer A_5 is subject to forces about which information is deemed to be primary and, additionally, the entire material is subject to forces which affect each of the layers. In this case, since the total force affecting the layer of constituent A_5 is different than the total forces affecting each of the other layers and information is deemed to be primary about the forces and resultant perturbation of the layer of constituent A_5 , attenuation terms due to constituents A_1 through A_4 and A_6 through A_N make up the secondary signal portion $n_{\lambda a}(t)$. Even if the additional forces which affect the entire material cause the same perturbation in each layer, including the layer of A_5 , the total forces on the layer of constituent A_5 cause it to have different total perturbation than each of the other layers of constituents A_1 through A_4 and A_6 through A_N .

It is often the case that the total perturbation affecting the layers associated with the secondary signal components is caused by random or erratic forces. This causes the thickness of layers to change erratically and the optical path length of each layer, $x_i(t)$, to change erratically, thereby producing a random or erratic secondary signal component $n_{\lambda a}(t)$. However, regardless of whether or not the secondary signal portion $n_{\lambda a}(t)$ is erratic, the secondary signal component $n_{\lambda a}(t)$ can be either removed or derived via a correlation canceler, such

as an adaptive noise canceler, having as one input, respectively, a secondary reference $n'(t)$ or a primary reference $s'(t)$ determined by a processor of the present invention as long as the perturbation on layers other than the layer of constituent A_5 is different than the perturbation on the layer of constituent A_5 . The correlation canceler yields a good approximation to either the primary signal $s_{\lambda a}(t)$ or the secondary signal $n_{\lambda a}(t)$. In the event that an approximation to the primary signal is obtained, the concentration of the constituent of interest, $c_5(t)$, can often be determined since in some physiological measurements, the thickness of the primary signal component, $x_5(t)$ in this example, is known or can be determined.

The correlation canceler utilized a sample of either the secondary reference $n'(t)$ or the primary reference $s'(t)$ determined from two substantially simultaneously measured signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$. $s_{\lambda a}(t)$ is determined as above in equation (7). $s_{\lambda b}(t)$ is determined similarly at a different wavelength λ_b . To find either the secondary reference $n'(t)$ or the primary reference $s'(t)$, attenuated transmitted energy is measured at the two different wavelengths λ_a and λ_b and transformed via logarithmic conversion. The signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ can then be written (logarithm converted) as:

$$s_{\lambda a}(t) = \varepsilon_5, \lambda_a c_5 x_5(t) + \sum_{i=1}^4 \varepsilon_i, \lambda_a c_i x_i + \sum_{i=6}^N \varepsilon_i, \lambda_a c_i x_i \quad (8)$$

$$s_{\lambda a}(t) = \varepsilon_5, \lambda_a c_5 x_5(t) + n_{\lambda a}(t) \quad (9)$$

$$s_{\lambda b}(t) = \varepsilon_5, \lambda_b c_5 x_5(t) + \sum_{i=1}^4 \varepsilon_i, \lambda_b c_i x_i + \sum_{i=6}^N \varepsilon_i, \lambda_b c_i x_i \quad (10)$$

$$s_{\lambda b}(t) = \varepsilon_5, \lambda_b c_5 x_5(t) + n_{\lambda b}(t) \quad (11)$$

Further transformations of the signals are the proportionality relationships defining ω_a and ω_v , similarly to equation (3), which allows determination of a noise reference $n'(t)$ and a primary reference $s'(t)$. These are:

$$\varepsilon_5, \lambda_a = \omega_a \varepsilon_5, \lambda_b \quad (12a)$$

$$n\lambda_a = \omega_v n\lambda_b \quad (12b)$$

where

$$n\lambda_a \neq \omega_a n\lambda_b \quad (13a)$$

$$\varepsilon_5, \lambda_a \neq \omega_v \varepsilon_5, \lambda_b. \quad (13b)$$

It is often the case that both equations (12) and (13) can be simultaneously satisfied. Multiplying equation (11) by ω_a and subtracting the result from equation (9) yields a non-zero secondary reference which is a linear sum of secondary signal components:

$$n'(t) = s_{\lambda_a}(t) - \omega_a s_{\lambda_b}(t) = n_{\lambda_a}(t) - \omega_a n_{\lambda_b}(t) \quad (14a)$$

$$= \sum_{i=1}^4 \varepsilon_{i,\lambda_a} c_i x_i(t) + \sum_{i=6}^N \varepsilon_{i,\lambda_a} c_i x_i(t) -$$

$$\sum_{i=1}^4 \omega_a \varepsilon_{i,\lambda_b} c_i x_i(t) + \sum_{i=6}^N \omega_a \varepsilon_{i,\lambda_b} c_i x_i(t) \quad (15a)$$

$$= \sum_{i=1}^4 c_i x_i(t) [\varepsilon_{i,\lambda_a} - \omega_a \varepsilon_{i,\lambda_b}] +$$

$$\sum_{i=6}^N c_i x_i(t) [\varepsilon_{i,\lambda_a} - \omega_a \varepsilon_{i,\lambda_b}] \quad (16a)$$

Multiplying equation (11) by ω_v and subtracting the

result from equation (9) yields a primary reference which is a linear sum of primary signal components:

$$s'(t) = s_{\lambda_a}(t) - \omega_v s_{\lambda_b}(t) = s_{\lambda_a}(t) - \omega_v s_{\lambda_b}(t) \quad (14b)$$

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$$= c_5 x_5(t) \varepsilon_5, \lambda_a - \omega_v c_5 x_5(t) \varepsilon_5, \lambda_b \quad (15b)$$

$$= c_5 x_5(t) [\varepsilon_5, \lambda_a - \omega_v \varepsilon_5, \lambda_b]. \quad (16b)$$

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A sample of either the secondary reference $n'(t)$ or the primary reference $s'(t)$, and a sample of either measured signal $s_{\lambda_a}(t)$ or $s_{\lambda_b}(t)$, are input to a correlation canceler 27, such as an adaptive noise canceler 30, an example of which is shown in Figures 5a and 5b and a preferred example of which is discussed herein under the heading PREFERRED, CORRELATION CANCELER USING A JOINT PROCESS ESTIMATOR IMPLEMENTATION. The correlation canceler 27 removes either the secondary portion $n_{\lambda_a}(t)$ or $n_{\lambda_b}(t)$, or the primary portions, $s_{\lambda_a}(t)$ or $s_{\lambda_b}(t)$, of the measured signal yielding a good approximation to either the primary signals $s''_{\lambda_a}(t) \approx \varepsilon_5, \lambda_a c_5 x_5(t)$ or $s''_{\lambda_b}(t) \approx \varepsilon_5, \lambda_b c_5 x_5(t)$ or the secondary signals $n''_{\lambda_a}(t) \approx n_{\lambda_a}(t)$ or $n''_{\lambda_b}(t) \approx n_{\lambda_b}(t)$. In the event that the primary signals are obtained, the concentration $c_5(t)$ may then be determined from the approximation to the primary signal $s''_{\lambda_a}(t)$ or $s''_{\lambda_b}(t)$ according to:

$$c_5(t) \approx s''_{\lambda_a}(t) / \varepsilon_5, \lambda_a x_5(t) \approx s''_{\lambda_b}(t) / \varepsilon_5, \lambda_b x_5(t). \quad (17)$$

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As discussed previously, the absorption coefficients are constant at each wavelength λ_a and λ_b and the thickness of the primary signal component, $x_5(t)$ in this example, is often known or can be determined as a function of time, thereby allowing calculation of the concentration $c_5(t)$ of

constituent A_5 .

DETERMINATION OF CONCENTRATION OR SATURATION

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IN A VOLUME CONTAINING MORE THAN ONE CONSTITUENT

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Referring to Figure 6b, another material having N different constituents arranged in layers is shown. In this material, two constituents A_5 and A_6 are found within one layer having thickness $x_{5,6}(t) = x_5(t) + x_6(t)$, located generally randomly within the layer. This is analogous to combining the layers of constituents A_5 and A_6 in Figure 6a. A combination of layers, such as the combination of layers of constituents A_5 and A_6 , is feasible when the two layers are under the same total forces which result in the same change of the optical path lengths $x_5(t)$ and $x_6(t)$ of the layers.

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Often it is desirable to find the concentration or the saturation, i.e., a percent concentration, of one constituent within a given thickness which contains more than one constituent and is subject to unique forces. A determination of the concentration or the saturation of a constituent within a given volume may be made with any number of constituents in the volume subject to the same total forces and therefore under the same perturbation or change. To determine the saturation of one constituent in a volume comprising many constituents, as many measured signals as there are constituents which absorb incident light energy are necessary. It will be understood that constituents which do not absorb light energy are not consequential in the determination of saturation. To determine the concentration, as many signals as there are constituents which absorb incident light energy are

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necessary as well as information about the sum of concentrations.

It is often the case that a thickness under unique motion contains only two constituents. For example, it may be desirable to know the concentration or saturation of A₅ within a given volume which contains A₅ and A₆. In this case, the primary signals s_{λa(t)} and s_{λb(t)} comprise terms related to both A₅ and A₆ so that a determination of the concentration or saturation of A₅ or A₆ in the volume may be made. A determination of saturation is discussed herein. It will be understood that the concentration of A₅ in a volume containing both A₅ and A₆ could also be determined if it is known that A₅+A₆=1, i.e., that there are no constituents in the volume which do not absorb incident light energy at the particular measurement wavelengths chosen. The measured signals s_{λa(t)} and s_{λb(t)} can be written (logarithm converted) as:

$$S_{\lambda a}(t) = \varepsilon_{5,\lambda a} c_{5,6}(t) + \varepsilon_{6,\lambda a} c_{6,5}(t) + n_{\lambda a}(t) \quad (18a)$$

$$= s_{\lambda a}(t) + n_{\lambda a}(t) \quad (18b)$$

$$S_{\lambda b}(t) = \varepsilon_{5,\lambda b} c_{5,6}(t) + \varepsilon_{6,\lambda b} c_{6,5}(t) + n_{\lambda b}(t) \quad (19a)$$

$$= s_{\lambda b}(t) + n_{\lambda b}(t). \quad (19b)$$

It is also often the case that there may be two or more thicknesses within a medium each containing the same two constituents but each experiencing a separate motion as in Figure 6c. For example, it may be desirable to know the concentration or saturation of A₅ within a given volume

which contains A_5 and A_6 as well as the concentration or
 saturation of A_3 within a given volume which contains A_3
 and A_4 , A_3 and A_4 having the same constituency as A_5 and
 A_6 , respectively. In this case, the primary signals $s_{\lambda a}(t)$
 and $s_{\lambda b}(t)$ again comprise terms related to both A_5 and A_6
 and portions of the secondary signals $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$
 comprise terms related to both A_3 and A_4 . The layers, A_3
 and A_4 , do not enter into the primary equation because they
 are assumed to be perturbed by random or erratic secondary
 forces which are uncorrelated with the primary force. Since
 constituents 3 and 5 as well as constituents 4 and 6 are
 taken to be the same, they have the same absorption
 coefficients. i. e. $\varepsilon_3, \lambda_a = \varepsilon_5, \lambda_a, \varepsilon_3, \lambda_b = \varepsilon_5, \lambda_b,$
 $\varepsilon_4, \lambda_a = \varepsilon_6, \lambda_a$ and $\varepsilon_4, \lambda_b = \varepsilon_6, \lambda_b$. Generally speaking,
 however, A_3 and A_4 will have different concentrations than
 A_5 and A_6 and will therefore have a different saturation.
 Consequently a single constituent within a medium may have
 one or more saturations associated with it. The primary and
 secondary signals according to this model may be written
 as:
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$$s_{\lambda a}(t) = [\varepsilon_5, \lambda_a c_5 + \varepsilon_6, \lambda_a c_6] x_5, 6(t) \quad (20a)$$

$$n_{\lambda a}(t) = [\varepsilon_5, \lambda_a c_3 + \varepsilon_6, \lambda_a c_4] x_3, 4(t)$$

$$+ \sum_{i=1}^2 \varepsilon_i, \lambda_a c_i x_i(t) + \sum_{i=7}^N \varepsilon_i, \lambda_a c_i x_i(t) \quad (20b)$$

$$n_{\lambda a}(t) = [\varepsilon_5, \lambda_a c_3 + \varepsilon_6, \lambda_a c_4] x_3, 4(t) + n_{\lambda a}(t) \quad (20c)$$

$$s_{\lambda b}(t) = [\varepsilon_5, \lambda_b c_5 + \varepsilon_6, \lambda_b c_6] x_5, 6(t) \quad (21a)$$

$$n_{\lambda b}(t) = [\varepsilon_5, \lambda_b c_3 + \varepsilon_6, \lambda_b c_4] x_3, 4(t)$$

$$+ \sum_{i=1}^2 \varepsilon_i, \lambda_b c_i x_i(t) + \sum_{i=7}^N \varepsilon_i, \lambda_b c_i x_i(t). \quad (21b)$$

$$n_{\lambda b}(t) = [\varepsilon_5, \lambda_b c_3 + \varepsilon_6, \lambda_b c_4] x_3, 4(t) + n_{\lambda b}(t) \quad (21c)$$

where signals $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ are similar to the secondary signals $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ except for the omission of the 3, 4 layer.

Any signal portions whether primary or secondary, outside of a known bandwidth of interest, including the constant undesired secondary signal portion resulting from the generally constant absorption of the constituents when not under perturbation, should be removed to determine an approximation to either the primary signal or the secondary signal within the bandwidth of interest. This is easily accomplished by traditional band pass filtering techniques. As in the previous example, it is often the case that the total perturbation or change affecting the layers associated with the secondary signal components is caused by random or erratic forces, causing the thickness of each layer, or the optical path length of each layer, $x_i(t)$, to change erratically, producing a random or erratic secondary signal component $n_{\lambda a}(t)$. Regardless of whether or not the secondary signal portion $n_{\lambda a}(t)$ is erratic, the secondary signal component $n_{\lambda a}(t)$ can be removed or derived via a correlation canceler, such as an adaptive noise canceler, having as one input a secondary reference $n'(t)$ or a primary reference $s'(t)$ determined by a processor of the present invention as long as the perturbation in layers other than the layer of constituents A_5 and A_6 is different than the perturbation in the layer of constituents A_5 and A_6 . Either the erratic secondary signal components $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ or the primary components $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ may advantageously be removed from equations (18) and (19), or alternatively equations (20) and (21), by a correlation canceler. The correlation canceler, again, requires a sample of either the primary reference $s'(t)$ or the secondary reference $n'(t)$ and a sample of either of the

composite signals $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ of equations (18) and (19).

DETERMINATION OF PRIMARY AND SECONDARY REFERENCE SIGNALS

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FOR SATURATION MEASUREMENTS

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Two methods which may be used by a processor of the present invention to determine either the secondary reference $n'(t)$ or the primary reference $s'(t)$ are a ratiometric method and a constant saturation method. One embodiment of a physiological monitor incorporating a processor of the present invention utilizes the ratiometric method wherein the two wavelengths λ_a and λ_b , at which the signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ are measured, are specifically chosen such that a relationship between the absorption coefficients $\varepsilon_{5,\lambda a}$, $\varepsilon_{5,\lambda b}$, $\varepsilon_{6,\lambda a}$ and $\varepsilon_{6,\lambda b}$ exists, i.e.:

$$\varepsilon_{5,\lambda a}/\varepsilon_{6,\lambda a} = \varepsilon_{5,\lambda b}/\varepsilon_{6,\lambda b} \quad (22)$$

The measured signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ can be factored and written as:

$$s_{\lambda a}(t) = \varepsilon_{6,\lambda a} [(\varepsilon_{5,\lambda a}/\varepsilon_{6,\lambda a}) c_5 \times_{5,6}(t) + c_6 \times_{5,6}(t)] + n_{\lambda a}(t) \quad (23a)$$

$$s_{\lambda a}(t) = \varepsilon_{6,\lambda a} [(\varepsilon_{5,\lambda a}/\varepsilon_{6,\lambda a}) c_5 \times_{5,6}(t) + c_6 \times_{5,6}(t) + (\varepsilon_{5,\lambda a}/\varepsilon_{6,\lambda a}) c_3 \times_{3,4}(t) + c_4 \times_{3,4}(t)] + n_{\lambda a}(t) \quad (23b)$$

$$s_{\lambda a}(t) = s_{\lambda a} + n_{\lambda a}(t) \quad (23c)$$

$$s_{\lambda b}(t) = \varepsilon_{6,\lambda b} [(\varepsilon_{5,\lambda b}/\varepsilon_{6,\lambda b}) c_5 \times_{5,6}(t) + c_6 \times_{5,6}(t)] + n_{\lambda b}(t) \quad (24a)$$

$$S_{\lambda b}(t) = \varepsilon_6, \lambda_b [(\varepsilon_5, \lambda_b / \varepsilon_6, \lambda_b) c_5 \times 5, 6(t) + c_6 \times 5, 6(t) \\ + (\varepsilon_5, \lambda_b / \varepsilon_6, \lambda_b) c_3 \times 3, 4(t) + c_4 \times 3, 4(t)] + n_{\lambda b}(t) \quad (24b)$$

$$S_{\lambda b}(t) = s_{\lambda b} + n_{\lambda b}(t). \quad (24c)$$

The wavelengths λ_a and λ_b , chosen to satisfy equation (22), cause the terms within the square brackets to be equal, thereby causing the terms other than $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ to be linearly dependent. Then, proportionality constants ω_{av} and ω_e may be found for the determination of a non-zero primary and secondary reference

$$\varepsilon_6, \lambda_a = \omega_{av} \varepsilon_6, \lambda_b \quad (25a)$$

$$n_{\lambda a}(t) = \omega_e n_{\lambda b}(t) \quad (25b)$$

$$\varepsilon_6, \lambda_a \neq \omega_e \varepsilon_6, \lambda_b \quad (26a)$$

$$n_{\lambda a}(t) \neq \omega_{av} n_{\lambda b}(t) \quad (26b)$$

It is often the case that both equations (25) and (26) can be simultaneously satisfied. Additionally, since the absorption coefficients of each constituent are constant with respect to wavelength, the proportionality constants ω_{av} and ω_e can be easily determined. Furthermore, absorption coefficients of other constituents A_1 through A_2 and A_7 through A_N are generally unequal to the absorption coefficients of A_3 , A_4 , A_5 and A_6 . Thus, the secondary components $n_{\lambda a}$ and $n_{\lambda b}$ are generally not made linearly dependent by the relationships of equations (22) and (25).

Multiplying equation (24) by ω_{av} and subtracting the

resulting equation from equation (23), a non-zero secondary reference is determined by:

$$n(t) = s_{\lambda_a}(t) - \omega_{av} s_{\lambda_b}(t)$$

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$$= s_{\lambda_a}(t) - \omega_{av} n_{\lambda_b}(t). \quad (27a)$$

Multiplying equation (24) by ω_e and subtracting the resulting equation from equation (23), a non-zero primary reference is determined by:

$$s(t) = s_{\lambda_a}(t) - \omega_e s_{\lambda_b}(t)$$

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$$= s_{\lambda_a}(t) - \omega_e s_{\lambda_b}(t). \quad (27b)$$

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An alternative method for determining reference signals from the measured signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ using a processor of the present invention is the constant saturation approach. In this approach, it is assumed that the saturation of A_5 in the volume containing A_5 and A_6 and the saturation of A_3 in the volume containing A_3 and A_4 remains relatively constant over some period of time, i.e.:

$$\text{Saturation}(A_5(t)) = c_5(t) / [c_5(t) + c_6(t)] \quad (28a)$$

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$$\text{Saturation}(A_3(t)) = c_3(t) / [c_3(t) + c_4(t)] \quad (28b)$$

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$$\text{Saturation}(A_5(t)) = \{1 + [c_6(t)/c_5(t)]\}^{-1} \quad (29a)$$

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$$\text{Saturation}(A_3(t)) = \{1 + [c_4(t)/c_3(t)]\}^{-1} \quad (29b)$$

are substantially constant over many samples of the measured signals s_{λ_a} and s_{λ_b} . This assumption is accurate over many samples since saturation generally changes

relatively slowly in physiological systems.

The constant saturation assumption is equivalent to assuming that:

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$$c_5(t)/c_6(t) = \text{constant}_1 \quad (30a)$$

$$c_3(t)/c_4(t) = \text{constant}_2 \quad (30b)$$

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since the only other term in equations (29a) and (29b) is a constant, namely the numeral 1.

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Using this assumption, the proportionality constants ω_a and ω_v which allow determination of the secondary reference signal $n'(t)$ and the primary reference signal $s'(t)$ in the constant saturation method are:

$$\omega_a = \frac{\varepsilon_5, \lambda_a c_5 x_5, 6(t) + \varepsilon_6, \lambda_a c_6 x_5, 6(t)}{\varepsilon_5, \lambda_b c_5 x_5, 6(t) + \varepsilon_6, \lambda_b c_6 x_5, 6(t)} \quad (31a)$$

$$= s\lambda_a(t)/s\lambda_b(t) \quad (32a)$$

$$= \frac{\varepsilon_5, \lambda_a c_5 + \varepsilon_6, \lambda_a c_6}{\varepsilon_5, \lambda_b c_5 + \varepsilon_6, \lambda_b c_6} \quad (33a)$$

$$= \frac{\varepsilon_5, \lambda_a (c_5/c_6) + \varepsilon_6, \lambda_a}{\varepsilon_5, \lambda_b (c_5/c_6) + \varepsilon_6, \lambda_b} \quad (34a)$$

$$\approx s''\lambda_a(t)/s''\lambda_b(t) = \text{constant}_3; \text{ where} \quad (35a)$$

$$n\lambda_a(t) \neq \omega_a(t)n\lambda_b(t) \quad (36a)$$

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and

$$\omega_v = \frac{\varepsilon_5, \lambda_a c_3 x_{3,4}(t) + \varepsilon_6, \lambda_a c_4 x_{3,4}(t)}{\varepsilon_5, \lambda_b c_3 x_{3,4}(t) + \varepsilon_6, \lambda_b c_4 x_{3,4}(t)} \quad (31b)$$

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$$= n\lambda_a(t)/n\lambda_b(t) \quad (32b)$$

$$= \frac{\varepsilon_5, \lambda_a c_3 + \varepsilon_6, \lambda_a c_4}{\varepsilon_5, \lambda_b c_3 + \varepsilon_6, \lambda_b c_4} \quad (33b)$$

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$$= \frac{\varepsilon_5, \lambda_a (c_3/c_4) + \varepsilon_6, \lambda_a}{\varepsilon_5, \lambda_b (c_3/c_4) + \varepsilon_6, \lambda_b} \quad (34b)$$

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$$\approx n''\lambda_a(t)/n''\lambda_b(t) = \text{constant}_4; \text{ where} \quad (35b)$$

$$s_{\lambda_a}(t) \neq \omega_v(t) s_{\lambda_b}(t). \quad (36b)$$

It is often the case that both equations (32) and (36) can be simultaneously satisfied to determine the proportionality constants ω_a and ω_v . Additionally, the absorption coefficients at each wavelength ε_5, λ_a , ε_6, λ_a , ε_5, λ_b , and ε_6, λ_b are constant and the central assumption of the constant saturation method is that $c_5(t)/c_6(t)$ and $c_3(t)/c_4(t)$ are constant over many sample periods. Thus, new proportionality constants ω_a and ω_v may be determined every few samples from new approximations to either the primary or secondary signal as output from the correlation canceler. Thus, the approximations to either the primary signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ or the secondary signals $n\lambda_a(t)$ and $n\lambda_b(t)$, found by the correlation canceler for a substantially immediately preceding set of samples of the measured signals $S_{\lambda_a}(t)$ and $S_{\lambda_b}(t)$ are used in a processor of the present invention for calculating the

proportionality constants, ω_a and ω_v , for the next set of samples of the measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$.

Multiplying equation (19) by ω_a and subtracting the resulting equation from equation (18) yields a non-zero secondary reference signal:

$$n'(t) = S_{\lambda a}(t) - \omega_a S_{\lambda b}(t) = n_{\lambda a}(t) - \omega_a n_{\lambda b}(t). \quad (37a)$$

Multiplying equation (19) by ω_v and subtracting the resulting equation from equation (18) yields a non-zero primary reference signal:

$$s'(t) = S_{\lambda a}(t) - \omega_v S_{\lambda b}(t) = s_{\lambda a}(t) - \omega_v s_{\lambda b}(t). \quad (37b)$$

When using the constant saturation method, it is not necessary for the patient to remain motionless for a short period of time such that an accurate initial saturation value can be determined by known methods other than correlation canceling. With no erratic, motion-induced signal portions, a physiological monitor can very quickly produce an initial value of the saturation of A_5 in the volume containing A_5 and A_6 . An example of a saturation calculation is given in the article "SPECTROPHOTOMETRIC DETERMINATION OF OXYGEN SATURATION OF BLOOD INDEPENDENT OF THE PRESENT OF INDOCYANINE GREEN" by G.A. Mook, et al., wherein determination of oxygen saturation in arterial blood is discussed. Another article discussing the calculation of oxygen saturation is "PULSE OXIMETRY: PHYSICAL PRINCIPLES, TECHNICAL REALIZATION AND PRESENT LIMITATIONS" by Michael R. Neuman. Then, with values for the coefficients ω_a and ω_v determined, a correlation canceler may be utilized with a secondary reference $n'(t)$ or a primary reference $s'(t)$ determined by the constant

saturation method.

DETERMINATION OF SIGNAL COEFFICIENTS FOR
PRIMARY AND SECONDARY REFERENCE SIGNALS
USING THE CONSTANT SATURATION METHOD

The reference processor 26 of Figures 4a and Figure 4b of the present invention may be configured to multiply the second measured signal $s_{\lambda b}(t) = s_{\lambda b}(t) + n_{\lambda b}(t)$ by a plurality of signal coefficients $\omega_1, \omega_2, \dots, \omega_n$ and then subtract each result from the first measured signal $s_{\lambda a}(t) = s_{\lambda a}(t) + n_{\lambda a}(t)$ to obtain a plurality of reference signals

$$r'(\omega, t) = s_{\lambda a}(t) - \omega s_{\lambda b}(t) + n_{\lambda a}(t) - \omega n_{\lambda b}(t) \quad (38)$$

for $\omega = \omega_1, \omega_2, \dots, \omega_n$ as shown in Figure 7a.

In order to determine either the primary reference $s'(t)$ or the secondary reference $n'(t)$ from the above plurality of reference signals of equation (38), signal coefficients ω_a and ω_v must be determined from the plurality of signal coefficients $\omega_1, \omega_2, \dots, \omega_n$. The coefficients ω_a and ω_v are such that they cause either the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ or the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ to cancel or nearly cancel when they are substituted into the reference function $r'(\omega, t)$, e. g.

$$s_{\lambda a}(t) = \omega_a s_{\lambda b}(t) \quad (39a)$$

$$n_{\lambda a}(t) = \omega_v n_{\lambda b}(t) \quad (39b)$$

$$n'(t) = r'(\omega_a, t) = n_{\lambda a}(t) - \omega_a n_{\lambda b}(t) \quad (39c)$$

$$s'(t) = r'(\omega_v, t) = s_{\lambda a}(t) - \omega_v s_{\lambda b}(t). \quad (39d)$$

In practice, one does not usually have significant prior information about either the primary signal portions

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$s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ or the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ of the measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$. The lack of this information makes it difficult to determine which of the plurality of coefficients $\omega_1, \omega_2, \dots, \omega_n$ correspond to the signal coefficients $\omega_a = s_{\lambda a}(t)/s_{\lambda b}(t)$ and $\omega_v = n_{\lambda a}(t)/n_{\lambda b}(t)$. Herein the preferred approach to determine the signal coefficients ω_a and ω_v from the plurality of coefficients $\omega_1, \omega_2, \dots, \omega_n$ employs the use of a correlation canceler 27, such as an adaptive noise canceler, which takes a first input which corresponds to one of the measured signals $S_{\lambda a}(t)$ or $S_{\lambda b}(t)$ and takes a second input which corresponds to successively each one of the plurality of reference signals $r'(\omega_1, t), r'(\omega_2, t), \dots, r'(\omega_n, t)$ as shown in Figure 7a. For each of the reference signals $r'(\omega_1, t), r'(\omega_2, t), \dots, r'(\omega_n, t)$ the corresponding output of the correlation canceler 27 is input to an integrator 29 for forming a cumulative output signal. The cumulative output signal is subsequently input to an extremum detector 31. The purpose of the extremum detector 31 is to chose signal coefficients ω_a and ω_v from the set $\omega_1, \omega_2, \dots, \omega_n$ by observing which provide a maximum in the cumulative output signal as in Figures 7b and 7c. In other words, coefficients which provide a maximum integrated output, such as energy or power, from the correlation canceler 27 correspond to the signal coefficients ω_a and ω_v . One could also configure a system geometry which would require one to locate the coefficients from the set $\omega_1, \omega_2, \dots, \omega_n$ which provide a minimum or inflection in the cumulative output signal to identify the signal coefficients ω_a and ω_v

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Use of a plurality of coefficients in the processor of the present invention in conjunction with a correlation canceler 27 to determine the signal coefficients ω_a and ω_v may be demonstrated by using the properties of correlation

cancellation. If x , y and z are taken to be any collection of three time varying signals, then the properties of a generic correlation canceler $C(x, y)$ may be defined as follows:

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Property (1) $C(x, y) = 0$ for x, y correlated

Property (2) $C(x, y) = x$ for x, y uncorrelated (40)

10 Property (3) $C(x + y, z) = C(x, z) + C(y, z).$

With properties (1), (2) and (3) it is easy to demonstrate that the energy or power output of a correlation canceler with a first input which corresponds to one of the measured signals $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ and a second input which corresponds to successively each one of a plurality of reference signals $r'(\omega_1, t)$, $r'(\omega_2, t)$, ..., $r'(\omega_n, t)$ can determine the signal coefficients ω_a and ω_b needed to produce the primary reference $s'(t)$ and secondary reference $n'(t)$. If we take as a first input to the correlation canceler the measured signal $s_{\lambda a}(t)$ and as a second input the plurality of reference signals $r'(\omega_1, t)$, $r'(\omega_2, t)$, ..., $r'(\omega_n, t)$ then the outputs of the correlation canceler $C(s_{\lambda a}(t), r'(\omega_j, t))$ for $j = 1, 2, \dots, n$ may be written as

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$$C(s_{\lambda a}(t) + n_{\lambda a}(t), s_{\lambda a}(t) - \omega_j s_{\lambda b}(t) + n_{\lambda a}(t) - \omega_j n_{\lambda b}(t)) \quad (41)$$

where $j = 1, 2, \dots, n$ and we have used the expressions

$$r'(\omega, t) = s_{\lambda a}(t) - \omega s_{\lambda b}(t) \quad (42)$$

$$s_{\lambda a}(t) = s_{\lambda a}(t) + n_{\lambda a}(t) \quad (43a)$$

$$s_{\lambda b}(t) = s_{\lambda b}(t) + n_{\lambda b}(t). \quad (43b)$$

The use of property (3) allows one to expand equation (41) into two terms

$$C(s_{\lambda a}(t), r'(\omega, t)) = C(s_{\lambda a}(t), s_{\lambda a}(t) - \omega s_{\lambda b}(t) + n_{\lambda a}(t) - \omega n_{\lambda b}(t))$$

$$(44) \quad + C(n_{\lambda_a}(t), s_{\lambda_a}(t) - \omega s_{\lambda_b}(t) + n_{\lambda_a}(t) - \omega n_{\lambda_b}(t))$$

so that upon use of properties (1) and (2) the correlation canceler output is given by

$$C(s_{\lambda_a}(t), r'(\omega_j, t)) = s_{\lambda_a}(t) \delta(\omega_j - \omega_a) + n_{\lambda_a}(t) \delta(\omega_j - \omega_v) \quad (45)$$

where $\delta(x)$ is the unit impulse function

$$\begin{aligned} \delta(x) &= 0 \text{ if } x \neq 0 \\ \delta(x) &= 1 \text{ if } x = 0. \end{aligned} \quad (46)$$

The time variable, t , of the correlation canceler output $C(s_{\lambda_a}(t), r'(\omega_j, t))$ may be eliminated by computing its energy or power. The energy of the correlation canceler output is given by

$$\begin{aligned} E_{\lambda_a}(\omega_j) &= \int C^2(s_{\lambda_a}(t), r'(\omega_j, t)) dt \\ &= \delta(\omega - \omega_a) \int s_{\lambda_a}^2(t) dt + \delta(\omega - \omega_v) \int n_{\lambda_a}^2(t) dt. \end{aligned} \quad (47a)$$

It must be understood that one could, equally well, have chosen the measured signal $s_{\lambda_b}(t)$ as the first input to the correlation canceler and the plurality of reference signals $r'(\omega_1, t), r'(\omega_2, t), \dots, r'(\omega_n, t)$ as the second input. In this event, the correlation canceler energy output is

$$\begin{aligned} E_{\lambda_b}(\omega) &= \int C^2(s_{\lambda_b}(t), r'(\omega, t)) dt \\ &= \delta(\omega - \omega_a) \int s_{\lambda_b}^2(t) dt + \delta(\omega - \omega_v) \int n_{\lambda_b}^2(t) dt. \end{aligned} \quad (47b)$$

It must also be understood that in practical situations the use of discrete time measurement signals may be employed as well as continuous time measurement signals. In the event that discrete time measurement signals are used integration approximation methods such as the trapezoid rule, midpoint

rule, Tick's rule, Simpson's approximation or other techniques may be used to compute the correlation canceler energy or power output. In the discrete time measurement signal case, the energy output of the correlation canceler may be written, using the trapezoid rule, as

$$E_{\lambda_a}(\omega) = \delta(\omega - \omega_a) \Delta t \{ \sum_{i=0}^n s^2 \lambda_a(t_i) - 0.5(s^2 \lambda_a(t_0) + s^2 \lambda_a(t_n)) \} + \delta(\omega - \omega_v) \Delta t \{ \sum_{i=0}^n n^2 \lambda_a(t_i) - 0.5(n^2 \lambda_a(t_0) + n^2 \lambda_a(t_n)) \} \quad (48a)$$

$$E_{\lambda_b}(\omega) = \delta(\omega - \omega_a) \Delta t \{ \sum_{i=0}^n s^2 \lambda_b(t_i) - 0.5(s^2 \lambda_b(t_0) + s^2 \lambda_b(t_n)) \} + \delta(\omega - \omega_v) \Delta t \{ \sum_{i=0}^n n^2 \lambda_b(t_i) - 0.5(n^2 \lambda_b(t_0) + n^2 \lambda_b(t_n)) \} \quad (48b)$$

where t_i is the i^{th} discrete time, t_0 is the initial time, t_n is the final time and Δt is the time between discrete time measurement samples.

The energy functions given above, and shown in Figure 7b, indicate that the correlation canceler output is usually zero due to correlation between the measured signal $s_{\lambda_a}(t)$ or $s_{\lambda_b}(t)$ and many of the plurality of reference signals $r'(\omega_1, t), r'(\omega_2, t), \dots, r'(\omega_n, t)r'(\omega, t)$. However, the energy functions are non zero at values of ω_j which correspond to cancellation of either the primary signal portions $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ or the secondary signal portions $n_{\lambda_a}(t)$ and $n_{\lambda_b}(t)$ in the reference signal $r'(\omega_j, t)$. These values correspond to the signal coefficients ω_a and ω_v .

It must be understood that there may be instances in time when either the primary signal portions $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ or the secondary signal portions $n_{\lambda_a}(t)$ and $n_{\lambda_b}(t)$ are identically zero or nearly zero. In these cases, only one signal coefficient value will provide maximum energy or power output of the correlation canceler.

Since there may be more than one signal coefficient value

which provides maximum correlation canceler energy or power output, an ambiguity may arise. It may not be immediately obvious which signal coefficient together with the reference function $r'(\omega, t)$ provides either the primary or secondary reference. In such cases, it is necessary to consider the constraints of the physical system at hand. For example, in pulse oximetry, it is known that arterial blood, whose signature is the primary plethysmographic wave, has greater oxygen saturation than venous blood, whose signature is the secondary erratic or random signal. Consequently, in pulse oximetry, the ratio of the primary signals due to arterial pulsation $\omega_a = s_{\lambda a}(t)/s_{\lambda b}(t)$ is the smaller of the two signal coefficient values while the ratio of the secondary signals due to mainly venous blood dynamics $\omega_v = n_{\lambda a}(t)/n_{\lambda b}(t)$ is the larger of the two signal coefficient values, assuming $\lambda a = 660$ nm and $\lambda b = 940$ nm.

It must be understood that in practical implementations of the plurality of reference signals and cross correlator technique, the ideal features listed as properties (1), (2) and (3) above will not be precisely satisfied but will be approximations thereof. Therefore, in practical implementations of the present invention, the correlation canceler energy curves depicted in Figure 7b will not consist of infinitely narrow delta functions but will have finite width associated with them as depicted in Figure 7c.

It should also be understood that it is possible to have more than two signal coefficient values which produce maximum energy or power output from a correlation canceler. This situation will arise when the measured signals each contain more than two components each of which are related by a ratio as follows:

$$S_{\lambda a}(t) = \sum_{i=1}^n f_{\lambda a, i}(t) \quad (49)$$

$$S_{\lambda b}(t) = \sum_{i=1}^n f_{\lambda b, i}(t)$$

5 where

$$f_{\lambda a, i}(t) = \omega_i f_{\lambda a, i}(t) \quad i = 1, \dots, n$$

$$\omega_i \neq \omega_j.$$

10 The ability to employ reference signal techniques together with a correlation cancellation, such as an adaptive noise canceler, to decompose a signal into two or more signal components each of which is related by a ratio is a further
15 aspect of the present invention.

PREFERRED CORRELATION CANCELER USING A

JOINT PROCESS ESTIMATOR IMPLEMENTATION

20 Once either the secondary reference $n'(t)$ or the primary reference $s'(t)$ is determined by the processor of the present invention using either the above described ratiometric or constant saturation methods, the correlation canceler can be implemented in either hardware or software.
25 The preferred implementation of a correlation canceler is that of an adaptive noise canceler using a joint process estimator.

30 The least mean squares (LMS) implementation of the internal processor 32 described above in conjunction with the adaptive noise canceler of Figures 5a and Figure 5b is relatively easy to implement, but lacks the speed of
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adaptation desirable for most physiological monitoring applications of the present invention. Thus, a faster approach for adaptive noise canceling, called a least-squares lattice joint process estimator model, is preferably used. A joint process estimator 60 is shown diagrammatically in Figure 8 and is described in detail in chapter 9 of Adaptive Filter Theory by Simon Haykin, published by Prentice-Hall, copyright 1986. This entire book, including Chapter 9, is hereby incorporated herein by reference. The function of the joint process estimator is to remove either the secondary signal portions $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$ or the primary signal portions $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ from the measured signals $S_{\lambda a}(t)$ or $S_{\lambda b}(t)$, yielding either a signal $s''_{\lambda a}(t)$ or $s''_{\lambda b}(t)$ or a signal $n''_{\lambda a}(t)$ or $n''_{\lambda b}(t)$ which is a good approximation to either the primary signal $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ or the secondary signal $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$. Thus, the joint process estimator estimates either the value of the primary signals $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ or the secondary signals $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$. The inputs to the joint process estimator 60 are either the secondary reference $n'(t)$ or the primary reference $s'(t)$ and the composite measured signal $S_{\lambda a}(t)$ or $S_{\lambda b}(t)$. The output is a good approximation to the signal $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ with either the secondary signal or the primary signal removed, i.e. a good approximation to either $s_{\lambda a}(t)$, $s_{\lambda b}(t)$, $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$.

The joint process estimator 60 of Figure 8 utilizes, in conjunction, a least square lattice predictor 70 and a regression filter 80. Either the secondary reference $n'(t)$ or the primary reference $s'(t)$ is input to the least square lattice predictor 70 while the measured signal $S_{\lambda a}(t)$ or $S_{\lambda b}(t)$ is input to the regression filter 80. For simplicity in the following description, $S_{\lambda a}(t)$ will be the measured

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signal from which either the primary portion $s_{\lambda a}(t)$ or the secondary portion $n_{\lambda a}(t)$ will be estimated by the joint process estimator 60. However, it will be noted that $s_{\lambda b}(t)$ could equally well be input to the regression filter 80 and the primary portion $s_{\lambda b}(t)$ or the secondary portion $n_{\lambda b}(t)$ of this signal could equally well be estimated.

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The joint process estimator 60 removes all frequencies that are present in both the reference $n'(t)$ or $s'(t)$, and the measured signal $S_{\lambda a}(t)$. The secondary signal portion $n_{\lambda a}(t)$ usually comprises frequencies unrelated to those of the primary signal portion $s_{\lambda a}(t)$. It is highly improbable that the secondary signal portion $n_{\lambda a}(t)$ would be of exactly the same spectral content as the primary signal portion $s_{\lambda a}(t)$. However, in the unlikely event that the spectral content of $s_{\lambda a}(t)$ and $n_{\lambda a}(t)$ are similar, this approach will not yield accurate results. Functionally, the joint process estimator 60 compares the reference input signal $n'(t)$ or $s'(t)$, which is correlated to either the secondary signal portion $n_{\lambda a}(t)$ or the primary signal portion $s_{\lambda a}(t)$, and input signal $S_{\lambda a}(t)$ and removes all frequencies which are identical. Thus, the joint process estimator 60 acts as a dynamic multiple notch filter to remove those frequencies in the secondary signal component $n_{\lambda a}(t)$ as they change erratically with the motion of the patient or those frequencies in the primary signal component $s_{\lambda a}(t)$ as they change with the arterial pulsation of the patient. This yields a signal having substantially the same spectral content and amplitude as either the primary signal $s_{\lambda a}(t)$ or the secondary signal $n_{\lambda a}(t)$. Thus, the output $s''_{\lambda a}(t)$ or $n''_{\lambda a}(t)$ of the joint process estimator 60 is a very good approximation to either the primary signal $s_{\lambda a}(t)$ or the secondary signal $n_{\lambda a}(t)$.

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The joint process estimator 60 can be divided into stages, beginning with a zero-stage and terminating in an m^{th} -stage, as shown in Figure 8. Each stage, except for the zero-stage, is identical to every other stage. The zero-stage is an input stage for the joint process estimator 60. The first stage through the m^{th} -stage work on the signal produced in the immediately previous stage, i.e., the $(m-1)^{\text{th}}$ -stage, such that a good approximation to either the primary signal $s''\lambda_a(t)$ or the secondary signal $n''\lambda_a(t)$ is produced as output from the m^{th} -stage.

The least-squares lattice predictor 70 comprises registers 90 and 92, summing elements 100 and 102, and delay elements 110. The registers 90 and 92 contain multiplicative values of a forward reflection coefficient $\Gamma_{f,m}(t)$ and a backward reflection coefficient $\Gamma_{b,m}(t)$ which multiply the reference signal $n'(t)$ or $s'(t)$ and signals derived from the reference signal $n'(t)$ or $s'(t)$. Each stage of the least-squares lattice predictor outputs a forward prediction error $f_m(t)$ and a backward prediction error $b_m(t)$. The subscript m is indicative of the stage.

For each set of samples, i.e. one sample of the reference signal $n'(t)$ or $s'(t)$ derived substantially simultaneously with one sample of the measured signal $s\lambda_a(t)$, the sample of the reference signal $n'(t)$ or $s'(t)$ is input to the least-squares lattice predictor 70. The zero-stage forward prediction error $f_0(t)$ and the zero-stage backward prediction error $b_0(t)$ are set equal to the reference signal $n'(t)$ or $s'(t)$. The backward prediction error $b_0(t)$ is delayed by one sample period by the delay element 110 in the first stage of the least-squares lattice predictor 70. Thus, the immediately previous value of the reference $n'(t)$ or $s'(t)$ is used in calculations involving

the first-stage delay element 110. The zero-stage forward prediction error is added to the negative of the delayed zero-stage backward prediction error $b_0(t-1)$ multiplied by the forward reflection coefficient value $\Gamma_{f,1}(t)$ register 90 value, to produce a first-stage forward prediction error $f_1(t)$. Additionally, the zero-stage forward prediction error $f_0(t)$ is multiplied by the backward reflection coefficient value $\Gamma_{b,1}(t)$ register 92 value and added to the delayed zero-stage backward prediction error $b_0(t-1)$ to produce a first-stage backward prediction error $b_1(t)$. In each subsequent stage, m , of the least square lattice predictor 70, the previous forward and backward prediction error values, $f_{m-1}(t)$ and $b_{m-1}(t-1)$, the backward prediction error being delayed by one sample period, are used to produce values of the forward and backward prediction errors for the present stage, $f_m(t)$ and $b_m(t)$.

The backward prediction error $b_m(t)$ is fed to the concurrent stage, m , of the regression filter 80. There it is input to a register 96, which contains a multiplicative regression coefficient value $\kappa_{m,\lambda_a}(t)$. For example, in the zero-stage of the regression filter 80, the zero-stage backward prediction error $b_0(t)$ is multiplied by the zero-stage regression coefficient $\kappa_{0,\lambda_a}(t)$ register 96 value and subtracted from the measured value of the signal $s_{\lambda_a}(t)$ at a summing element 106 to produce a first stage estimation error signal $e_{1,\lambda_a}(t)$. The first-stage estimation error signal $e_{1,\lambda_a}(t)$ is a first approximation to either the primary signal or the secondary signal. This first-stage estimation error signal $e_{1,\lambda_a}(t)$ is input to the first-stage of the regression filter 80. The first-stage backward prediction error $b_1(t)$, multiplied by the first-stage regression coefficient $\kappa_{1,\lambda_a}(t)$ register 96 value is subtracted from the first-stage estimation error signal

$e_{1,\lambda a}(t)$ to produce the second-stage estimation error $e_{2,\lambda a}(t)$. The second-stage estimation error signal $e_{2,\lambda a}(t)$ is a second, somewhat better approximation to either the primary signal $s_{\lambda a}(t)$ or the secondary signal $n_{\lambda a}(t)$.

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The same processes are repeated in the least-squares lattice predictor 70 and the regression filter 80 for each stage until a good approximation $e_{m,\lambda a}(t)$, to either the primary signal $s_{\lambda a}(t)$ or the secondary signal $n_{\lambda a}(t)$ is determined. Each of the signals discussed above, including the forward prediction error $f_m(t)$, the backward prediction error $b_m(t)$, the estimation error signal $e_{m,\lambda a}(t)$, is necessary to calculate the forward reflection coefficient $\Gamma_{f,m}(t)$, the backward reflection coefficient $\Gamma_{b,m}(t)$, and the regression coefficient $\kappa_{m,\lambda a}(t)$ register 90, 92, and 96 values in each stage, m . In addition to the forward prediction error $f_m(t)$, the backward prediction error $b_m(t)$, and the estimation error $e_{m,\lambda a}(t)$ signals, a number of intermediate variables, not shown in Figure 8 but based on the values labeled in Figure 8, are required to calculate the forward reflection coefficient $\Gamma_{f,m}(t)$, the backward reflection coefficient $\Gamma_{b,m}(t)$, and the regression coefficient $\kappa_{m,\lambda a}(t)$ register 90, 92, and 96 values.

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Intermediate variables include a weighted sum of the forward prediction error squares $\bar{\gamma}_m(t)$, a weighted sum of the backward prediction error squares $\beta_m(t)$, a scalar parameter $\Delta_m(t)$, a conversion factor $\gamma_m(t)$, and another scalar parameter $\rho_{m,\lambda a}(t)$. The weighted sum of the forward prediction errors $\bar{\gamma}_m(t)$ is defined as:

$$\mathfrak{J}_m(t) = \sum_{i=1}^t \lambda^{t-i} |f_m(i)|^2; \quad (50)$$

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where λ without a wavelength identifier, a or b, is a constant multiplicative value unrelated to wavelength and is typically less than or equal to one, i.e., $\lambda \leq 1$. The weighted sum of the backward prediction errors $\beta_m(t)$ is defined as:

$$\beta_m(t) = \sum_{i=1}^t \lambda^{t-i} |b_m(i)|^2 \quad (51)$$

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where, again, λ without a wavelength identifier, a or b, is a constant multiplicative value unrelated to wavelength and is typically less than or equal to one, i.e., $\lambda \leq 1$. These weighted sum intermediate error signals can be manipulated such that they are more easily solved for, as described in Chapter 9, § 9.3. and defined hereinafter in equations (65) and (66).

DESCRIPTION OF THE JOINT PROCESS ESTIMATOR

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The operation of the joint process estimator 60 is as follows. When the joint process estimator 60 is turned on, the initial values of intermediate variables and signals including the parameter $\Delta_{m-1}(t)$, the weighted sum of the forward prediction error signals $\mathfrak{J}_{m-1}(t)$, the weighted sum of the backward prediction error signals $\beta_{m-1}(t)$, the parameter $\rho_m, \lambda_a(t)$, and the zero-stage estimation error $e_0, \lambda_a(t)$ are initialized, some to zero and some to a small positive number δ :

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5 (52)

$$\Delta_{m-1}(0) = 0;$$

10 (53)

$$\tilde{\gamma}_{m-1}(0) = \delta;$$

15 (54)

$$\beta_{m-1}(0) = \delta;$$

20 (55)

$$\rho_{m,\lambda a}(0) = 0;$$

25 (56)

$$e_{0,\lambda a}(t) = s_{\lambda a}(t) \text{ for } t \geq 0.$$

After initialization, a simultaneous sample of the measured signal $s_{\lambda a}(t)$ or $s_{\lambda b}(t)$ and either the secondary reference $n'(t)$ or the primary reference $s'(t)$ are input to the joint process estimator 60, as shown in Figure 8. The forward and backward prediction error signals $f_0(t)$ and $b_0(t)$, and intermediate variables including the weighted sums of the forward and backward error signals $\tilde{\gamma}_0(t)$ and $\beta_0(t)$, and the conversion factor $\gamma_0(t)$ are calculated for the zero-stage according to:

25 (57a)

$$f_0(t) = b_0(t) = n'(t)$$

30 (58a)

$$\tilde{\gamma}_0(t) = \beta_0(t) = \lambda \tilde{\gamma}_0(t-1) + |n'(t)|^2$$

35 (59a)

$$\gamma_0(t-1) = 1$$

if a secondary reference $n'(t)$ is used or according to:

30 (57b)

$$f_0(t) = b_0(t) = s'(t)$$

35 (58b)

$$\tilde{\gamma}_0(t) = \beta_0(t) = \lambda \tilde{\gamma}_0(t-1) + |s'(t)|^2$$

40 (59b)

$$\gamma_0(t-1) = 1$$

if a primary reference $s'(t)$ is used where, again, λ without a wavelength identifier, a or b, is a constant multiplicative value unrelated to wavelength.

5 Forward reflection coefficient $\Gamma_{f,m}(t)$, backward reflection coefficient $\Gamma_{b,m}(t)$, and regression coefficient $\kappa_m, \lambda_a(t)$ register 90, 92 and 96 values in each stage thereafter are set according to the output of the previous stage. The forward reflection coefficient $\Gamma_{f,1}(t)$, backward reflection coefficient $\Gamma_{b,1}(t)$, and regression coefficient $\kappa_1, \lambda_a(t)$ register 90, 92 and 96 values in the first stage 10 are thus set according to algorithm using values in the zero-stage of the joint process estimator 60. In each stage, $m \geq 1$, intermediate values and register values including the parameter $\Delta_{m-1}(t)$; the forward reflection coefficient $\Gamma_{f,m}(t)$ register 90 value; the backward reflection coefficient $\Gamma_{b,m}(t)$ register 92 value; the forward and backward error signals $f_m(t)$ and $b_m(t)$; the weighted sum of squared forward prediction errors $\mathfrak{J}_{f,m}(t)$, as manipulated in § 9.3 of the Haykin book; the weighted sum of squared backward prediction errors $\beta_{b,m}(t)$, as manipulated in § 9.3 of the Haykin book; the conversion factor $\gamma_m(t)$; the parameter $\rho_m, \lambda_a(t)$; the regression coefficient $\kappa_m, \lambda_a(t)$ register 96 value; and the estimation error $e_{m+1}\lambda_a(t)$ value are set according to:

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$$\Delta_{m-1}(t) = \lambda \Delta_{m-1}(t-1) + \{b_{m-1}(t-1)f^*_{m-1}(t)/\gamma_{m-1}(t-1)\} \quad (60)$$

$$\Gamma_{f,m}(t) = -\{\Delta_{m-1}(t)/\beta_{m-1}(t-1)\} \quad (61)$$

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$$\Gamma_{b,m}(t) = -\{\Delta^*_{m-1}(t)/\mathfrak{J}_{m-1}(t)\} \quad (62)$$

$$f_m(t) = f_{m-1}(t) + \Gamma^*_{f,m}(t)b_{m-1}(t-1) \quad (63)$$

$$b_m(t) = b_{m-1}(t-1) + \Gamma^*_{b,m}(t) f_{m-1}(t) \quad (64)$$

$$\tilde{\gamma}_m(t) = \tilde{\gamma}_{m-1}(t) - \{ |\Delta_{m-1}(t)|^2 / \beta_{m-1}(t-1) \} \quad (65)$$

$$\beta_m(t) = \beta_{m-1}(t-1) - \{ |\Delta_{m-1}(t)|^2 / \tilde{\gamma}_{m-1}(t) \} \quad (66)$$

$$\gamma_m(t-1) = \gamma_{m-1}(t-1) - \{ |b_{m-1}(t-1)|^2 / \beta_{m-1}(t-1) \} \quad (67)$$

$$\rho_{m,\lambda a}(t) = \lambda \rho_{m,\lambda a}(t-1) + \{ b_m(t) e^*_{m,\lambda a}(t) / \gamma_m(t) \} \quad (68)$$

$$\kappa_{m,\lambda a}(t) = \{ \rho_{m,\lambda a}(t) / \beta_m(t) \} \quad (69)$$

$$e_{m+1,\lambda a}(t) = e_{m,\lambda a}(t) - \kappa^*_{m,\lambda a}(t) b_m(t) \quad (70)$$

where a (*) denotes a complex conjugate.

These equations cause the error signals $f_m(t)$, $b_m(t)$, $e_{m,\lambda a}(t)$ to be squared or to be multiplied by one another, in effect squaring the errors, and creating new intermediate error values, such as $\Delta_{m-1}(t)$. The error signals and the intermediate error values are recursively tied together, as shown in the above equations (60) through (70). They interact to minimize the error signals in the next stage.

After a good approximation to either the primary signal $s_{\lambda a}(t)$ or the secondary signal $n_{\lambda a}(t)$ has been determined by the joint process estimator 60, a next set of samples, including a sample of the measured signal $s_{\lambda a}(t)$ and a sample of either the secondary reference $n'(t)$ or the primary reference $s'(t)$, are input to the joint process estimator 60. The re-initialization process does not re-occur, such that the forward and backward reflection coefficient $\Gamma_{f,m}(t)$ and $\Gamma_{b,m}(t)$ register 90, 92 values and

the regression coefficient $\kappa_m, \lambda_a(t)$ register 96 value reflect the multiplicative values required to estimate either the primary signal portion $s_{\lambda_a}(t)$ or the secondary signal portion $n_{\lambda_a}(t)$ of the sample of $s_{\lambda_a}(t)$ input previously. Thus, information from previous samples is used to estimate either the primary or secondary signal portion of a present set of samples in each stage.

FLOWCHART OF JOINT PROCESS ESTIMATOR

In a signal processor, such as a physiological monitor, incorporating a reference processor of the present invention to determine a reference $n'(t)$ or $s'(t)$ for input to a correlation canceler, a joint process estimator 60 type adaptive noise canceler is generally implemented via a software program having an iterative loop. One iteration of the loop is analogous to a single stage of the joint process estimator as shown in Figure 8. Thus, if a loop is iterated m times, it is equivalent to an m stage joint process estimator 60.

A flow chart of a subroutine to estimate the primary signal portion $s_{\lambda_a}(t)$ or the secondary signal portion $n_{\lambda_a}(t)$ of a measured signal, $S_{\lambda_a}(t)$ is shown in Figure 9. The flow chart describes how the action of a reference processor for determining either the secondary reference or the primary reference and the joint process estimator 60 would be implemented in software.

A one-time only initialization is performed when the physiological monitor is turned on, as indicated by an "INITIALIZE NOISE CANCELER" box 120. The initialization sets all registers 90, 92, and 96 and delay element variables 110 to the values described above in equations

(52) through (56).

5 Next, a set of simultaneous samples of the measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ is input to the subroutine represented by the flowchart in Figure 9. Then a time update of each of the delay element program variables occurs, as indicated in a "TIME UPDATE OF $[Z^{-1}]$ ELEMENTS" box 130, wherein the value stored in each of the delay element variables 110 is set to the value at the input of the delay element variable 110. Thus, the zero-stage backward prediction error $b_0(t)$ is stored in the first-stage delay element variable, the first-stage backward prediction error $b_1(t)$ is stored in the second-stage delay element variable, and so on.

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30 Then, using the set of measured signal samples $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$, the reference signal is calculated according to the ratiometric or the constant saturation method described above. This is indicated by a "CALCULATE REFERENCE $[n'(t)$ or $s'(t)]$ FOR TWO MEASURED SIGNAL SAMPLES" box 140.

A zero-stage order update is performed next as indicated in a "ZERO-STAGE UPDATE" box 150. The zero-stage backward prediction error $b_0(t)$, and the zero-stage forward prediction error $f_0(t)$ are set equal to the value of the reference signal $n'(t)$ or $s'(t)$. Additionally, the weighted sum of the forward prediction errors $\beta_m(t)$ and the weighted sum of backward prediction errors $\beta_m(t)$ are set equal to the value defined in equations (53) and (54).

35 Next, a loop counter, m , is initialized as indicated in a " $m=0$ " box 160. A maximum value of m , defining the total number of stages to be used by the subroutine corresponding to the flowchart in Figure 9, is also defined. Typically, the loop is constructed such that it

stops iterating once a criterion for convergence upon a best approximation to either the primary signal or the secondary signal has been met by the joint process estimator 60. Additionally, a maximum number of loop iterations may be chosen at which the loop stops iteration. In a preferred embodiment of a physiological monitor of the present invention, a maximum number of iterations, $m=6$ to $m=10$, is advantageously chosen.

Within the loop, the forward and backward reflection coefficient $\Gamma_{f,m}(t)$ and $\Gamma_{b,m}(t)$ register 90 and 92 values in the least-squares lattice filter are calculated first, as indicated by the "ORDER UPDATE MTH CELL OF LSL-LATTICE" box 170 in Figure 9. This requires calculation of intermediate variable and signal values used in determining register 90, 92, and 96 values in the present stage, the next stage, and in the regression filter 80.

The calculation of regression filter register 96 value $\kappa_m, \lambda_a(t)$ is performed next, indicated by the "ORDER UPDATE MTH STAGE OF REGRESSION FILTER(S)" box 180. The two order update boxes 170 and 180 are performed in sequence m times, until m has reached its predetermined maximum (in the preferred embodiment, $m=6$ to $m=10$) or a solution has been converged upon, as indicated by a YES path from a "DONE" decision box 190. In a computer subroutine, convergence is determined by checking if the weighted sums of the forward and backward prediction errors $\beta_m(t)$ and $\beta_m(t)$ are less than a small positive number. An output is calculated next, as indicated by a "CALCULATE OUTPUT" box 200. The output is a good approximation to either the primary signal or secondary signal, as determined by the reference processor 26 and joint process estimator 60 subroutine corresponding to the flow chart of Figure 9. This is displayed (or used

in a calculation in another subroutine), as indicated by a "TO DISPLAY" box 210.

5 A new set of samples of the two measured signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ is input to the processor and joint process estimator 60 adaptive noise canceler subroutine corresponding to the flowchart of Figure 9 and the process reiterates for these samples. Note, however, that the initialization process does not re-occur. New sets of
10 measured signal samples $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ are continuously input to the reference processor 26 and joint process estimator 60 adaptive noise canceler subroutine. The output forms a chain of samples which is representative of a continuous wave. This waveform is a good approximation to either the primary signal waveform $s_{\lambda a}(t)$ or the secondary waveform $n_{\lambda a}(t)$ at wavelength λa . The waveform may also be a good approximation to either the primary signal waveform $s_{\lambda b}(t)$ or the secondary waveform $n''_{\lambda b}(t)$ at wavelength λb .

CALCULATION OF SATURATION FROM

CORRELATION CANCELER OUTPUT

25 Physiological monitors may use the approximation of the primary signals $s''_{\lambda a}(t)$ or $s''_{\lambda b}(t)$ or the secondary signals $n''_{\lambda a}(t)$ or $n''_{\lambda b}(t)$ to calculate another quantity, such as the saturation of one constituent in a volume containing that constituent plus one or more other constituents. Generally, such calculations require information about either a primary or secondary signal at
30 two wavelengths. For example, the constant saturation method requires a good approximation of the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ of both measured signals $s_{\lambda a}(t)$

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and $s_{\lambda b}(t)$. Then, the arterial saturation is determined from the approximations to both signals, i.e. $s''_{\lambda a}(t)$ and $s''_{\lambda b}(t)$. The constant saturation method also requires a good approximation of the secondary signal portions $n_{\lambda a}(t)$ or $n_{\lambda b}(t)$. Then an estimate of the venous saturation may be determined from the approximations to these signals i. e. $n''_{\lambda a}(t)$ and $n''_{\lambda b}(t)$.

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In other physiological measurements, information about a signal at a third wavelength is necessary. For example, to find the saturation using the ratiometric method, signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ are used to find the reference signal $n'(t)$ or $s'(t)$. But as discussed previously, λ_a and λ_b were chosen to satisfy a proportionality relationship like that of equation (22). This proportionality relationship forces the two primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ of equations (23c) and (24c) to be linearly dependent. Generally, linearly dependent mathematical equations cannot be solved for the unknowns. Analogously, some desirable information cannot be derived from two linearly dependent signals. Thus, to determine the saturation using the ratiometric method, a third signal is simultaneously measured at wavelength λ_c . The wavelength λ_c is chosen such that the primary portion $s_{\lambda c}(t)$ of the measured signal $S_{\lambda c}(t)$ is not linearly dependent with the primary portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ of the measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$. Since all measurements are taken substantially simultaneously, the secondary reference signal $n'(t)$ is correlated to the secondary signal portions $n_{\lambda a}$, $n_{\lambda b}$, and $n_{\lambda c}$ of each of the measured signals $S_{\lambda a}(t)$, $S_{\lambda b}(t)$, and $S_{\lambda c}(t)$ and can be used to estimate approximations to the primary signal portions $s_{\lambda a}(t)$, $s_{\lambda b}(t)$, and $s_{\lambda c}(t)$ for all three measured signals $S_{\lambda a}(t)$, $S_{\lambda b}(t)$, and $S_{\lambda c}(t)$. Using the ratiometric method, estimation

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of the ratio of signal portions $s_{\lambda a}(t)$ and $s_{\lambda c}(t)$ of the two measured signals $S_{\lambda a}(t)$ and $S_{\lambda c}(t)$, chosen correctly, is usually satisfactory to determine most physiological data.

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A joint process estimator 60 having two regression filters 80a and 80b is shown in Figure 10. A first regression filter 80a accepts a measured signal $S_{\lambda a}(t)$. A second regression filter 80b accepts a measured signal $S_{\lambda b}(t)$ or $S_{\lambda c}(t)$, depending whether the constant saturation method or the ratiometric method is used to determine the reference signal $n'(t)$ or $s'(t)$ for the constant saturation method or $n'(t)$ or $s'(t)$ for the ratiometric method. The first and second regression filters 80a and 80b are independent. The backward prediction error $b_m(t)$ is input to each regression filter 80a and 80b, the input for the second regression filter 80b bypassing the first regression filter 80a.

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The second regression filter 80b comprises registers 98, and summing elements 108 arranged similarly to those in the first regression filter 80a. The second regression filter 80b operates via an additional intermediate variable in conjunction with those defined by equations (60) through (70), i.e.:

$$\rho_{m,\lambda b}(t) = \lambda \rho_{m,\lambda b}(t-1) + \{b_m(t)e^{*}_{m,\lambda b}(t)/\gamma_m(t)\}; \text{ or } (71)$$

$$\rho_{m,\lambda c}(t) = \lambda \rho_{m,\lambda c}(t-1) + \{b_m(t)e^{*}_{m,\lambda c}(t)/\gamma_m(t)\}; \text{ and } (72)$$

$$\rho_{0,\lambda b}(0) = 0; \text{ or } (73)$$

$$\rho_{0,\lambda c}(0) = 0. \text{ (74)}$$

The second regression filter 80b has an error signal value defined similar to the first regression filter error signal values, $e_{m+1,\lambda a}(t)$, i.e.:

$$e_{m+1, \lambda b}(t) = e_{m, \lambda b}(t) - \kappa^*_{m, \lambda b}(t)b_m(t); \text{ or} \quad (75)$$

$$e_{m+1}, \lambda_C(t) = e_m, \lambda_C(t) - \kappa^*_{m, \lambda_B(t)} b_m(t); \text{ and} \quad (76)$$

$$e_{0,\lambda b}(t) = s_{\lambda b}(t) \text{ for } t \geq 0; \text{ or} \quad (77)$$

$$e_{0,\lambda_C}(t) = s_{\lambda_C}(t) \text{ for } t \geq 0. \quad (78)$$

The second regression filter has a regression coefficient $\kappa_m, \lambda_b(t)$ register 98 value defined similarly to the first regression filter error signal values, i.e.:

$$\kappa_m, \lambda_b(t) = \{\rho_m, \lambda_b(t)/\beta_m(t)\}; \text{ or} \quad (79)$$

$$\kappa_m, \lambda_C(t) = \{\rho_m, \lambda_C(t)/\beta_m(t)\}; \quad (80)$$

These values are used in conjunction with those intermediate variable values, signal values, register and register values defined in equations (52) through (70). These signals are calculated in an order defined by placing the additional signals immediately adjacent a similar signal for the wavelength λ_a .

For the ratiometric method, $S_{\lambda C}(t)$ is input to the second regression filter 80b. The output of the second regression filter 80b is then a good approximation to the primary signal $s''\lambda_C(t)$ or secondary signal $n''\lambda_C(t)$. For the constant saturation method, $S_{\lambda B}(t)$ is input to the second regression filter 80b. The output is then a good

approximation to the primary signal $s''\lambda_b(t)$ or secondary signal $s''\lambda_b(t)$.

The addition of the second regression filter 80b does not substantially change the computer program subroutine represented by the flowchart of Figure 9. Instead of an order update of the m^{th} stage of only one regression filter, an order update of the m^{th} stage of both regression filters 80a and 80b is performed. This is characterized by the plural designation in the "ORDER UPDATE OF m^{th} STAGE OF REGRESSION FILTER(S)" box 180 in Figure 9. Since the regression filters 80a and 80b operate independently, independent calculations can be performed in the reference processor and joint process estimator 60 adaptive noise canceler subroutine modeled by the flowchart of Figure 9.

CALCULATION OF SATURATION

Once good approximations to the primary signal portions, $s''\lambda_a(t)$ and $s''\lambda_c(t)$ or the secondary signal portions $n''\lambda_a(t)$ and $n''\lambda_c(t)$ for the ratiometric method and $s''\lambda_a(t)$ and $s''\lambda_b(t)$ or $n''\lambda_a(t)$ and $n''\lambda_c(t)$ for the constant saturation method, have been determined by the joint process estimator 60, the saturation of A_5 in a volume containing A_5 and A_6 , for example, may be calculated according to various known methods. Mathematically, the approximations to the primary signals can be written:

$$\begin{aligned} s''\lambda_a(t) &\approx \varepsilon_5, \lambda_a c_{5,6}(t) + \varepsilon_6, \lambda_a c_{6,5}(t) \\ &+ \varepsilon_5, \lambda_a c_{3,4}(t) + \varepsilon_6, \lambda_a c_{4,3}(t) \end{aligned} \quad (81a)$$

$$s''\lambda_c(t) \approx \varepsilon_5, \lambda_c c_5 x_{5,6}(t) + \varepsilon_6, \lambda_c c_6 x_{5,6}(t) \\ + \varepsilon_5, \lambda_c c_3 x_{3,4}(t) + \varepsilon_6, \lambda_c c_4 x_{3,4}(t) \quad (82a)$$

for the ratiometric method using wavelengths λ_a and λ_c , and assuming that the secondary reference $n'(t)$ is uncorrelated with $x_{3,4}(t)$ and $x_{5,6}(t)$. Terms involving $x_{3,4}(t)$ and $x_{5,6}(t)$ may then be separated using the constant saturation method. It is important to understand that if $n'(t)$ is uncorrelated with $x_{3,4}(t)$ and $x_{5,6}(t)$, use of the ratiometric method followed by use of the constant saturation method results in a more accurate computation of the saturation of A_3 in the layer $x_{3,4}$ than by use of the ratiometric or constant saturation methods alone. In the event that $n'(t)$ and $x_{3,4}(t)$ are correlated the ratiometric method yields

$$s''\lambda_a(t) \approx \varepsilon_5, \lambda_a c_5 x_{5,6}(t) + \varepsilon_6, \lambda_a c_6 x_{5,6}(t); \text{ and} \quad (81b)$$

$$s''\lambda_c(t) \approx \varepsilon_5, \lambda_c c_5 x_{5,6}(t) + \varepsilon_6, \lambda_c c_6 x_{5,6}(t). \quad (82b)$$

For the constant saturation method, the approximations to the primary signals can be written, in terms of λ_a and λ_b , as:

$$s''\lambda_a(t) \approx \varepsilon_5, \lambda_a c_5 x_{5,6}(t) + \varepsilon_6, \lambda_a c_6 x_{5,6}(t); \text{ and} \quad (83)$$

$$s''\lambda_b(t) \approx \varepsilon_5, \lambda_b c_5 x_{5,6}(t) + \varepsilon_6, \lambda_b c_6 x_{5,6}(t). \quad (84)$$

Equations (81b), (82b), (83) and (84) are equivalent to two equations having three unknowns, namely $c_5(t)$, $c_6(t)$ and $x_{5,6}(t)$. In both the ratiometric and the constant saturation cases, the saturation can be determined by acquiring approximations to the primary or secondary signal

portions at two different, yet proximate times t_1 and t_2 over which the saturation of A_5 in the volume containing A_5 and A_6 and the saturation of A_3 in the volume containing A_3 and A_4 does not change substantially. For example, for the primary signals estimated by the ratiometric method, at times t_1 and t_2 :

$$s''\lambda_a(t_1) \approx \varepsilon_5, \lambda_a c_{5,6}(t_1) + \varepsilon_6, \lambda_a c_{6,5}(t_1) \quad (85)$$

$$s''\lambda_c(t_1) \approx \varepsilon_5, \lambda_c c_{5,6}(t_1) + \varepsilon_6, \lambda_c c_{6,5}(t_1) \quad (86)$$

$$s''\lambda_a(t_2) \approx \varepsilon_5, \lambda_a c_{5,6}(t_2) + \varepsilon_6, \lambda_a c_{6,5}(t_2) \quad (87)$$

$$s''\lambda_c(t_2) \approx \varepsilon_5, \lambda_c c_{5,6}(t_2) + \varepsilon_6, \lambda_c c_{6,5}(t_2) \quad (88)$$

Then, difference signals may be determined which relate the signals of equations (85) through (88), i.e.:

$$\Delta s_{\lambda_a} = s''\lambda_a(t_1) - s''\lambda_a(t_2) \approx \varepsilon_5, \lambda_a c_{5,6} \Delta x + \varepsilon_6, \lambda_a c_{6,5} \Delta x; \text{ and } \quad (89)$$

$$\Delta s_{\lambda_c} = s''\lambda_c(t_1) - s''\lambda_c(t_2) \approx \varepsilon_5, \lambda_c c_{5,6} \Delta x + \varepsilon_6, \lambda_c c_{6,5} \Delta x; \quad (90)$$

where $\Delta x = x_{5,6}(t_1) - x_{5,6}(t_2)$. The average saturation at time $t = (t_1 + t_2)/2$ is:

$$\text{Saturation}(t) = c_5(t) / [c_5(t) + c_6(t)] \quad (91)$$

$$= \frac{\varepsilon_6, \lambda_a - \varepsilon_6, \lambda_c (\Delta s_{\lambda_a} / \Delta s_{\lambda_c})}{\varepsilon_6, \lambda_a - \varepsilon_5, \lambda_a - (\varepsilon_6, \lambda_c - \varepsilon_5, \lambda_c) (\Delta s_{\lambda_a} / \Delta s_{\lambda_c})} \quad (92)$$

It will be understood that the Δx term drops out from the saturation calculation because of the division. Thus, knowledge of the thickness of the primary constituents is not required to calculate saturation.

PULSE OXIMETRY MEASUREMENTS

A specific example of a physiological monitor utilizing a processor of the present invention to determine a secondary reference $n'(t)$ for input to a correlation canceler that removes erratic motion-induced secondary signal portions is a pulse oximeter. Pulse oximetry may also be performed utilizing a processor of the present invention to determine a primary signal reference $s'(t)$ which may be used for display purposes or for input to a correlation canceler to derive information about patient movement and venous blood oxygen saturation.

A pulse oximeter typically causes energy to propagate through a medium where blood flows close to the surface for example, an ear lobe, or a digit such as a finger, or a forehead. An attenuated signal is measured after propagation through or reflected from the medium. The pulse oximeter estimates the saturation of oxygenated blood.

Freshly oxygenated blood is pumped at high pressure from the heart into the arteries for use by the body. The volume of blood in the arteries varies with the heartbeat, giving rise to a variation in absorption of energy at the rate of the heartbeat, or the pulse.

Oxygen depleted, or deoxygenated, blood is returned to the heart by the veins along with unused oxygenated blood. The volume of blood in the veins varies with the rate of breathing, which is typically much slower than the heartbeat. Thus, when there is no motion induced variation in the thickness of the veins, venous blood causes a low frequency variation in absorption of energy. When there is

motion induced variation in the thickness of the veins, the low frequency variation in absorption is coupled with the erratic variation in absorption due to motion artifact.

5 In absorption measurements using the transmission of energy through a medium, two light emitting diodes (LED's) are positioned on one side of a portion of the body where blood flows close to the surface, such as a finger, and a photodetector is positioned on the opposite side of the finger. Typically, in pulse oximetry measurements, one LED emits a visible wavelength, preferably red, and the other LED emits an infrared wavelength. However, one skilled in the art will realize that other wavelength combinations could be used.

10 The finger comprises skin, tissue, muscle, both arterial blood and venous blood, fat, etc., each of which absorbs light energy differently due to different absorption coefficients, different concentrations, and different thicknesses. When the patient is not moving, absorption is substantially constant except for the flow of blood. The constant attenuation can be determined and subtracted from the signal via traditional filtering techniques. When the patient moves, the absorption becomes erratic. Erratic motion induced noise typically cannot be predetermined and/or subtracted from the measured signal via traditional filtering techniques. Thus, determining the oxygen saturation of arterial blood and venous blood becomes more difficult.

15 A schematic of a physiological monitor for pulse oximetry is shown in Figure 11. Two LED's 300 and 302, one LED 300 emitting red wavelengths and another LED 302 emitting infrared wavelengths, are placed adjacent a finger

310. A photodetector 320, which produces an electrical signal corresponding to the attenuated visible and infrared light energy signals is located opposite the LED's 300 and 302. The photodetector 320 is connected to a single channel of common processing circuitry including an amplifier 330 which is in turn connected to a band pass filter 340. The band pass filter 340 passes its output signal into a synchronized demodulator 350 which has a plurality of output channels. One output channel is for signals corresponding to visible wavelengths and another output channel is for signals corresponding to infrared wavelengths.

The output channels of the synchronized demodulator for signals corresponding to both the visible and infrared wavelengths are each connected to separate paths, each path comprising further processing circuitry. Each path includes a DC offset removal element 360 and 362, such as a differential amplifier, a programmable gain amplifier 370 and 372 and a low pass filter 380 and 382. The output of each low pass filter 380 and 382 is amplified in a second programmable gain amplifier 390 and 392 and then input to a multiplexer 400.

The multiplexer 400 is connected to an analog-to-digital converter 410 which is in turn connected to a microprocessor 420. Control lines between the microprocessor 420 and the multiplexer 400, the microprocessor 420 and the analog-to-digital converter 410, and the microprocessor 420 and each programmable gain amplifier 370, 372, 390, and 392 are formed. The microprocessor 420 has additional control lines, one of which leads to a display 430 and the other of which leads to an LED driver 440 situated in a feedback loop with the

two LED's 300 and 302.

The LED's 300 and 302 each emits energy which is absorbed by the finger 310 and received by the photodetector 320. The photodetector 320 produces an electrical signal which corresponds to the intensity of the light energy striking the photodetector 320 surface. The amplifier 330 amplifies this electrical signal for ease of processing. The band pass filter 340 then removes unwanted high and low frequencies. The synchronized demodulator 350 separates the electrical signal into electrical signals corresponding to the red and infrared light energy components. A predetermined reference voltage, V_{ref} , is subtracted by the DC offset removal element 360 and 362 from each of the separate signals to remove substantially constant absorption which corresponds to absorption when there is no motion induced signal component. Then the first programmable gain amplifiers 370 and 372 amplify each signal for ease of manipulation. The low pass filters 380 and 382 integrate each signal to remove unwanted high frequency components and the second programmable gain amplifiers 390 and 392 amplify each signal for further ease of processing.

The multiplexer 400 acts as an analog switch between the electrical signals corresponding to the red and the infrared light energy, allowing first a signal corresponding to the red light to enter the analog-to-digital converter 410 and then a signal corresponding to the infrared light to enter the analog-to-digital converter 410. This eliminates the need for multiple analog-to-digital converters 410. The analog-to-digital converter 410 inputs the data into the microprocessor 420 for calculation of either a primary or secondary reference signal via the

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processing technique of the present invention and removal or derivation of motion induced signal portions via a correlation canceler, such as an adaptive noise canceler. The microprocessor 420 centrally controls the multiplexer 400, the analog-to-digital converter 410, and the first and second programmable gain amplifiers 370 and 390 for both the red and the infrared channels. Additionally, the microprocessor 420 controls the intensity of the LED's 302 and 304 through the LED driver 440 in a servo loop to keep the average intensity received at the photodetector 320 within an appropriate range. Within the microprocessor 420 a reference signal $n'(t)$ or $s'(t)$ is calculated via either the constant saturation method or the ratiometric method, as described above, the constant saturation method being generally preferred. This signal is used in an adaptive noise canceler of the joint process estimator type 60, as described above.

The multiplexer 400 time multiplexes, or sequentially switches between, the electrical signals corresponding to the red and the infrared light energy. This allows a single channel to be used to detect and begin processing the electrical signals. For example, the red LED 300 is energized first and the attenuated signal is measured at the photodetector 320. An electrical signal corresponding to the intensity of the attenuated red light energy is passed to the common processing circuitry. The infrared LED 302 is energized next and the attenuated signal is measured at the photodetector 320. An electrical signal corresponding to the intensity of the attenuated infrared light energy is passed to the common processing circuitry. Then, the red LED 300 is energized again and the corresponding electrical signal is passed to the common processing circuitry. The sequential energization of LED's

300 and 302 occurs continuously while the pulse oximeter is operating.

The processing circuitry is divided into distinct paths after the synchronized demodulator 350 to ease time constraints generated by time multiplexing. In the preferred embodiment of the pulse oximeter shown in Figure 11, a sample rate, or LED energization rate, of .625 Hz is advantageously employed. Thus, electrical signals reach the synchronized demodulator 350 at a rate of 625 Hz. Time multiplexing is not used in place of the separate paths due to settling time constraints of the low pass filters 380, 382, and 384.

In Figure 11, a third LED 304 is shown adjacent the finger, located near the LED's 300 and 302. The third LED 304 is used to measure a third signal $S_{\lambda c}(t)$ to be used to determine saturation using the ratiometric method. The third LED 304 is time multiplexed with the red and infrared LED's 300 and 302. Thus, a third signal is input to the common processing circuitry in sequence with the signals from the red and infrared LED's 300 and 302. After passing through and being processed by the operational amplifier 330, the band pass filter 340, and the synchronized demodulator 350, the third electrical signal corresponding to light energy at wavelength λc is input to a separate path including a DC offset removal element 364, a first programmable gain amplifier 374, a low pass filter 384, and a second programmable gain amplifier 394. The third signal is then input to the multiplexer 400.

The dashed line connection for the third LED 304 indicates that this third LED 304 is incorporated into the pulse oximeter when the ratiometric method is used; it is

unnecessary for the constant saturation method. When the third LED 304 is used, the multiplexer 400 acts as an analog switch between all three LED 300, 302, and 304 signals. If the third LED 304 is utilized, feedback loops between the microprocessor 420 and the first and second programmable gain amplifier 374 and 394 in the λ_c wavelength path are also formed.

For pulse oximetry measurements using the ratiometric method, the signals (logarithm converted) transmitted through the finger 310 at each wavelength λ_a , λ_b , and λ_c are:

$$S_{\lambda_a}(t) = S_{\lambda \text{red}1}(t) = \epsilon_{HbO_2, \lambda_a} c^A_{HbO_2} x^A(t) + \epsilon_{Hb, \lambda_a} c^A_{Hb} x^A(t) + \epsilon_{HbO_2, \lambda_a} c^V_{HbO_2} x^V(t) + \epsilon_{Hb, \lambda_a} c^V_{Hb} x^V(t) + n_{\lambda_a}(t). \quad (93)$$

$$S_{\lambda_b}(t) = S_{\lambda \text{red}2}(t) = \epsilon_{HbO_2, \lambda_b} c^A_{HbO_2} x^A(t) + \epsilon_{Hb, \lambda_b} c^A_{Hb} x^A(t) + \epsilon_{HbO_2, \lambda_b} c^V_{HbO_2} x^V(t) + \epsilon_{Hb, \lambda_b} c^V_{Hb} x^V(t) + n_{\lambda_b}(t). \quad (94)$$

$$S_{\lambda_c}(t) = S_{\lambda \text{IR}}(t) = \epsilon_{HbO_2, \lambda_c} c^A_{HbO_2} x^A(t) + \epsilon_{Hb, \lambda_c} c^A_{Hb} x^A(t) + \epsilon_{HbO_2, \lambda_c} c^V_{HbO_2} x^V(t) + \epsilon_{Hb, \lambda_c} c^V_{Hb} x^V(t) + n_{\lambda_c}(t). \quad (95)$$

In equations (93) through (95), $x^A(t)$ is the lump-sum thickness of the arterial blood in the finger; $x^V(t)$ is the lump-sum thickness of venous blood in the finger; $\epsilon_{HbO_2, \lambda_a}$, $\epsilon_{HbO_2, \lambda_b}$, $\epsilon_{HbO_2, \lambda_c}$, ϵ_{Hb, λ_a} , ϵ_{Hb, λ_b} , and ϵ_{Hb, λ_c} are the absorption coefficients of the oxygenated and non-oxygenated hemoglobin, at each wavelength measured; and $c_{HbO_2}(t)$ and $c_{Hb}(t)$ with the superscript designations A and V are the concentrations of the oxygenated and non-oxygenated arterial blood and venous blood, respectively.

For the ratiometric method, the wavelengths chosen are typically two in the visible red range, i.e., λ_a and λ_b , and one in the infrared range, i.e., λ_c . As described above, the measurement wavelengths λ_a and λ_b are advantageously chosen to satisfy a proportionality relationship which removes the primary signal portions $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$, yielding a secondary reference $n'(t)$. In the preferred embodiment, the ratiometric method is used to determine the secondary reference signal $n'(t)$ by picking two wavelengths that cause the primary portions $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ of the measured signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ to become linearly dependent similarly to equation (22); i.e. wavelengths λ_a and λ_b which satisfy:

$$\varepsilon_{Hb02,\lambda_a}/\varepsilon_{Hb,\lambda_a} = \varepsilon_{Hb02,\lambda_b}/\varepsilon_{Hb,\lambda_b} \quad (96)$$

Typical wavelength values chosen are $\lambda_a = 650\text{nm}$ and $\lambda_b = 685\text{nm}$. Additionally a typical wavelength value for λ_c is $\lambda_c = 940\text{nm}$. By picking wavelengths λ_a and λ_b to satisfy equation (96) the venous portion of the measured signal is also caused to become linearly dependent even though it is not usually considered to be part of the primary signals as is the case in the constant saturation method. Thus, the venous portion of the signal is removed with the primary portion of the constant saturation method. The proportionality relationship between equations (93) and (94) which allows determination of a non-zero secondary reference signal $n'(t)$, similarly to equation (25) is:

$$\omega_{av} = \varepsilon_{Hb,\lambda_a}/\varepsilon_{Hb,\lambda_b}; \text{ where} \quad (97)$$

$$n_{\lambda_a}(t) \neq \omega_{av} n_{\lambda_b}(t). \quad (98)$$

In pulse oximetry, both equations (97) and (98) can typically be satisfied simultaneously.

Figure 12 is a graph of the absorption coefficients of oxygenated and deoxygenated hemoglobin (ϵ_{HbO_2} and ϵ_{Hb}) vs. wavelength (λ). Figure 13 is a graph of the ratio of the absorption coefficients vs. wavelength, i.e., $\epsilon_{Hb}/\epsilon_{HbO_2}$ vs. λ over the range of wavelength within circle 13 in Figure 12. Anywhere a horizontal line touches the curve of Figure 13 twice, as does line 400, the condition of equation (96) is satisfied. Figure 14 shows an exploded view of the area of Figure 12 within the circle 13. Values of ϵ_{HbO_2} and ϵ_{Hb} at the wavelengths where a horizontal line touches the curve of Figure 13 twice can then be determined from the data in Figure 14 to solve for the proportionality relationship of equation (97).

A special case of the ratiometric method is when the absorption coefficients ϵ_{HbO_2} and ϵ_{Hb} are equal at a wavelength. Arrow 410 in Figure 12 indicates one such location, called an isobestic point. Figure 14 shows an exploded view of the isobestic point. To use isobestic points with the ratiometric method, two wavelengths at isobestic points are determined to satisfy equation (96)

Multiplying equation (94) by ω_{av} and then subtracting equation (94) from equation (93), a non-zero secondary reference signal $n'(t)$ is determined by:

$$n'(t) = s_{\lambda_a}(t) - \omega_{av}s_{\lambda_b}(t) = n_{\lambda_a}(t) - \omega_{av}n_{\lambda_b}. \quad (99)$$

This secondary reference signal $n'(t)$ has spectral content corresponding to the erratic, motion-induced noise. When it is input to a correlation canceler, such as an adaptive noise canceler, with either the signals $s_{\lambda_a}(t)$ and

5 $s_{\lambda c}(t)$ or $s_{\lambda b}(t)$ and $s_{\lambda c}(t)$ input to two regression filters
 10 80a and 80b as in Figure 10, the adaptive noise canceler
 will function much like an adaptive multiple notch filter
 and remove frequency components present in both the
 secondary reference signal $n'(t)$ and the measured signals
 from the measured signals $s_{\lambda a}(t)$ and $s_{\lambda c}(t)$ or $s_{\lambda b}(t)$ and
 $s_{\lambda c}(t)$. If the secondary reference signal $n'(t)$ is
 correlated to the venous portion, then the adaptive noise
 canceler is able to remove erratic noise caused in the
 venous portion of the measured signals $s_{\lambda a}(t)$, $s_{\lambda b}(t)$, and
 $s_{\lambda c}(t)$ even though the venous portion of the measured
 signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ was not incorporated in the
 secondary reference signal $n'(t)$. In the event that the
 secondary reference signal $n'(t)$ is not correlated to the
 venous component, then, the adaptive noise canceler
 generally will not remove the venous portion from the
 measured signals. However, a band pass filter applied to
 the approximations to the primary signals $s''_{\lambda a}(t)$ and
 $s''_{\lambda c}(t)$ or $s''_{\lambda b}(t)$ and $s''_{\lambda c}(t)$ can remove the low frequency
 20 venous signal due to breathing.

25 For pulse oximetry measurements using the constant
 saturation method, the signals (logarithm converted)
 transmitted through the finger 310 at each wavelength λ_a
 and λ_b are:

$$s_{\lambda a}(t) = s_{\lambda red1}(t) = \varepsilon_{HbO_2, \lambda_a} c^A_{HbO_2} x^A(t) + \varepsilon_{Hb, \lambda_a} c^A_{Hb} x^A(t) + \varepsilon_{HbO_2, \lambda_a} c^V_{HbO_2} x^V(t) + \varepsilon_{Hb, \lambda_a} c^V_{Hb} x^V(t) + n_{\lambda a}(t) \quad (100a)$$

$$s_{\lambda a}(t) = \varepsilon_{HbO_2, \lambda_a} c^A_{HbO_2} x^A(t) + \varepsilon_{Hb, \lambda_a} c^A_{Hb} x^A(t) + n_{\lambda a}(t) \quad (100b)$$

$$s_{\lambda a}(t) = s_{\lambda a}(t) + n_{\lambda a}(t) \quad (100c)$$

$$S_{\lambda b}(t) = S_{\lambda red2}(t) = \varepsilon_{HbO_2, \lambda b} c^A_{HbO_2} x^A(t) + \varepsilon_{Hb, \lambda b} c^A_{Hb} x^A(t) \\ (101a)$$

$$+ \varepsilon_{HbO_2, \lambda b} c^V_{HbO_2} x^V(t) + \varepsilon_{Hb, \lambda b} c^V_{Hb} x^V(t) + n_{\lambda b}(t)$$

5 $S_{\lambda b}(t) = \varepsilon_{HbO_2, \lambda b} c^A_{HbO_2} x^A(t) + \varepsilon_{Hb, \lambda b} c^A_{Hb} x^A(t) + n_{\lambda b}(t) \quad (101b)$

10 $s_{\lambda b}(t) = s_{\lambda b}(t) + n_{\lambda b}(t) \quad (101c)$

15 For the constant saturation method, the wavelengths chosen are typically one in the visible red range, i.e., λ_a , and one in the infrared range, i.e., λ_b . Typical wavelength values chosen are $\lambda_a = 660\text{nm}$ and $\lambda_b = 940\text{nm}$. Using the constant saturation method, it is assumed that $c^A_{HbO_2}(t)/c^A_{Hb}(t) = \text{constant}_1$ and $c^V_{HbO_2}(t)/c^V_{Hb}(t) = \text{constant}_2$. The oxygen saturation of arterial and venous blood changes slowly, if at all, with respect to the sample rate, making this a valid assumption. The proportionality factors for equations (100) and (101) can then be written as:

20 $\omega_a(t) = \frac{\varepsilon_{HbO_2, \lambda_a} c^A_{HbO_2} x(t) + \varepsilon_{Hb, \lambda_a} c^A_{Hb} x(t)}{\varepsilon_{HbO_2, \lambda_b} c^A_{HbO_2} x(t) + \varepsilon_{Hb, \lambda_b} c^A_{Hb} x(t)} \quad (102)$

25 $s_{\lambda a}(t) = \omega_a(t) s_{\lambda b}(t) \quad (103a)$

30 $n_{\lambda a}(t) \neq \omega_a(t) n_{\lambda b}(t) \quad (104a)$

$n_{\lambda a}(t) = \omega_v(t) n_{\lambda b}(t) \quad (103b)$

$s_{\lambda a}(t) \neq \omega_v(t) s_{\lambda b}(t) \quad (104b)$

35 In pulse oximetry, it is typically the case that both

equations (103) and (104) can be satisfied simultaneously.

Multiplying equation (101) by $\omega_a(t)$ and then subtracting equation (101) from equation (100), a non-zero secondary reference signal $n'(t)$ is determined by:

$$n'(t) = s_{\lambda a}(t) - \omega_a(t)s_{\lambda b}(t) \quad (105a)$$

$$\begin{aligned} &= \varepsilon_{HbO_2, \lambda a} c^V_{HbO_2} x^V(t) + \varepsilon_{Hb, \lambda a} c^V_{Hb} x^V(t) + n_{\lambda a}(t) \\ &\quad - \omega_a(t) [\varepsilon_{HbO_2, \lambda b} c^V_{HbO_2} x^V(t) + \varepsilon_{Hb, \lambda b} c^V_{Hb} x^V(t) + n_{\lambda b}(t)]. \end{aligned} \quad (106a)$$

Multiplying equation (101) by $\omega_v(t)$ and then subtracting equation (101) from equation (100), a non-zero primary reference signal $s'(t)$ is determined by:

$$s'(t) = s_{\lambda a}(t) - \omega_v(t)s_{\lambda b}(t) \quad (105b)$$

$$= s_{\lambda a}(t) - \omega_v(t)s_{\lambda b}(t) \quad (106b)$$

The constant saturation assumption does not cause the venous contribution to the absorption to be canceled along with the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$, as did the relationship of equation (96) used in the ratiometric method. Thus, frequencies associated with both the low frequency modulated absorption due to venous absorption when the patient is still and the erratically modulated absorption due to venous absorption when the patient is moving are represented in the secondary reference signal $n'(t)$. Thus, the correlation canceler can remove or derive both erratically modulated absorption due to venous blood in the finger under motion and the constant low frequency cyclic absorption of venous blood.

Using either method, a primary reference $s'(t)$ or a secondary reference $n'(t)$ is determined by the processor of the present invention for use in a correlation canceler, such as an adaptive noise canceler, which is defined by software in the microprocessor. The preferred adaptive noise canceler is the joint process estimator 60 described above.

Illustrating the operation of the ratiometric method of the present invention, Figures 15, 16 and 17 show signals measured for use in determining the saturation of oxygenated arterial blood using a reference processor of the present invention which employs the ratiometric method, i.e., the signals $S_{\lambda a}(t) = S_{\lambda red1}(t)$, $S_{\lambda b}(t) = S_{\lambda red2}(t)$, and $S_{\lambda c}(t) = S_{\lambda IR}(t)$. A first segment 15a, 16a, and 17a of each of the signals, is relatively undisturbed by motion artifact, i.e., the patient did not move substantially during the time period in which these segments were measured. These segments 15a, 16a, and 17a are thus generally representative of the plethysmographic waveform at each of the measured wavelengths. These waveforms are taken to be the primary signals $s_{\lambda a}(t)$, $s_{\lambda b}(t)$, and $s_{\lambda c}(t)$. A second segment 15b, 16b, and 17b of each of the signals is affected by motion artifact, i.e., the patient did move during the time period in which these segments were measured. Each of these segments 15b, 16b, and 17b shows large motion induced excursions in the measured signal. These waveforms contain both primary plethysmographic signals and secondary motion induced excursions. A third segment 15c, 16c, and 17c of each of the signals is again relatively unaffected by motion artifact and is thus generally representative of the plethysmographic waveform at each of the measured wavelengths.

Figure 18 shows the secondary reference signal $n'(t) = n_{\lambda a} - \omega_{av}n_{\lambda b}(t)$, as determined by a reference processor of the present invention utilizing the ratiometric method. As discussed previously, the secondary reference signal $n'(t)$ is correlated to the secondary signal portions $n_{\lambda a}$, $n_{\lambda b}$, and $n_{\lambda c}$. Thus, a first segment 18a of the secondary reference signal $n'(t)$ is generally flat, corresponding to the fact that there is very little motion induced noise in the first segments 15a, 16a, and 17a of each signal. A second segment 18b of the secondary reference signal $n'(t)$ exhibits large excursions, corresponding to the large motion induced excursions in each of the measured signals. A third segment 18c of the secondary reference signal $n'(t)$ is generally flat, again corresponding to the lack of motion artifact in the third segments 15c, 16c, and 17c of each measured signal.

Figure 19 shows the primary reference signal $s'(t) = s_{\lambda a} - \omega_{es}\lambda_b(t)$, as determined by a reference processor of the present invention utilizing the ratiometric method. As discussed previously, the primary reference signal $s'(t)$ is correlated to the primary signal portions $s_{\lambda a}(t)$, $s_{\lambda b}(t)$, and $s_{\lambda c}(t)$. Thus, a first segment 19a of the primary reference signal $s'(t)$ is indicative of the plethysmographic waveform, corresponding to the fact that there is very little motion induced noise in the first segments 15a, 16a, and 17a of each signal. A second segment 19b of the primary reference signal $s'(t)$ also exhibits a signal related to a plethysmographic waveform, corresponding to each of the measured signals in the absence of the large motion induced excursions. A third segment 19c of the primary reference signal $s'(t)$ is generally indicative of the plethysmographic waveform, again corresponding to the

lack of motion artifact in the third segments 15c, 16c, and 17c of each measured signal.

Figures 20 and 21 show the approximations $s''\lambda_a(t)$ and $s''\lambda_c(t)$ to the primary signals $s_{\lambda_a}(t)$ and $s_{\lambda_c}(t)$ as estimated by the correlation canceler 27 using a secondary reference signal $n'(t)$ determined by the ratiometric method. Figures 20 and 21 illustrate the effect of correlation cancelation using the secondary reference signal $n'(t)$ as determined by the reference processor of the present invention using the ratiometric method. Segments 20b and 21b are not dominated by motion induced noise as were segments 15b, 16b, and 17b of the measured signals. Additionally, segments 20a, 21a, 20c, and 21c have not been substantially changed from the measured signal segments 15a, 17a, 15c, and 17c where there was no motion induced noise.

Figures 22 and 23 show the approximations $n''\lambda_a(t)$ and $n''\lambda_c(t)$ to the primary signals $n_{\lambda_a}(t)$ and $n_{\lambda_c}(t)$ as estimated by the correlation canceler 27 using a primary reference signal $s'(t)$ determined by the ratiometric method. Note that the scale of Figures 15 through 23 is not the same for each figure to better illustrate changes in each signal. Figures 22 and 23 illustrate the effect of correlation cancelation using the primary reference signal $s'(t)$ as determined by the reference processor of the present invention using the ratiometric method. Only segments 22b and 23b are dominated by motion induced noise as were segments 15b, 16b, and 17b of the measured signals. Additionally, segments 22a, 23a, 22c, and 23c are nearly zero corresponding to the measured signal segments 15a, 17a, 15c, and 17c where there was no motion induced noise.

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5 Illustrating the operation of the constant saturation
10 method of the present invention, Figures 24 and 25 show
15 signals measured for input to a reference processor of the
20 present invention which employs the constant saturation
25 method, i.e., the signals $s_{\lambda a}(t) = s_{\lambda red}(t)$ and
30 $s_{\lambda b}(t) = s_{\lambda IR}(t)$. A first segment 24a and 25a of each of
35 the signals is relatively undisturbed by motion artifact,
40 i.e., the patient did not move substantially during the
45 time period in which these segments were measured. These
50 segments 24a and 25a are thus generally representative of
55 the primary plethysmographic waveform at each of the
60 measured wavelengths. A second segment 24b and 25b of each
65 of the signals is affected by motion artifact, i.e., the
70 patient did move during the time period in which these
75 segments were measured. Each of these segments 24b and 25b
80 shows large motion induced excursions in the measured
85 signal. A third segment 24c and 25c of each of the signals
90 is again relatively unaffected by motion artifact and is
95 thus generally representative of the primary
wavelengths.

5 Figure 26 shows the secondary reference signal
10 $n'(t) = n_{\lambda a}(t) - \omega_a n_{\lambda b}(t)$, as determined by a reference
15 processor of the present invention utilizing the constant
20 saturation method. Again, the secondary reference signal
25 portion $n'(t)$ is correlated to the secondary signal portions
30 $n_{\lambda a}$ and $n_{\lambda b}$. Thus, a first segment 26a of the secondary
35 reference signal $n'(t)$ is generally flat, corresponding to
40 the fact that there is very little motion induced noise in
45 the first segments 24a and 25a of each signal. A second
50 segment 26b of the secondary reference signal $n'(t)$
55 exhibits large excursions, corresponding to the large
60 motion induced excursions in each of the measured signals.

A third segment 26c of the noise reference signal $n'(t)$ is generally flat, again corresponding to the lack of motion artifact in the third segments 24c and 25c of each measured signal.

, Figure 27 shows the primary reference signal $s'(t) = s_{\lambda a} - \omega_v s_{\lambda b}(t)$, as determined by a reference processor of the present invention utilizing the constant saturation method. As discussed previously, the primary reference signal $s'(t)$ is correlated to the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$. Thus, a first segment 27a of the primary reference signal $s'(t)$ is indicative of the plethysmographic waveform, corresponding to the fact that there is very little motion induced noise in the first segments 24a and 25a of each signal. A second segment 27b of the primary reference signal $s'(t)$ also exhibits a signal related to a plethysmographic waveform, corresponding to each of the measured signals in the absence of the large motion induced excursions. A third segment 27c of the primary reference signal $s'(t)$ is generally indicative of the plethysmographic waveform, again corresponding to the lack of motion artifact in the third segments 24c and 25c of each measured signal.

Figures 28 and 29 show the approximations $s''\lambda_a(t)$ and $s''\lambda_b(t)$ to the primary signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ as estimated by the correlation canceler 27 using a secondary reference signal $n'(t)$ determined by the constant saturation method. Figures 28 and 29 illustrate the effect of correlation cancellation using the secondary reference signal $n'(t)$ as determined by a reference processor of the present invention utilizing the constant saturation method. Segments 28b and 28b are not dominated by motion induced noise as were segments 24b and 25b of the measured signals.

Additionally, segments 28a, 29a, 28c, and 29c have not been substantially changed from the measured signal segments 24a, 25a, 24c, and 25c where there was no motion induced noise.

Figures 30 and 31 show the approximations $n''\lambda_a(t)$ and $n''\lambda_b(t)$ to the secondary signals $n\lambda_a(t)$ and $n\lambda_b(t)$ as estimated by the correlation canceler 27 using a primary reference signal $s'(t)$ determined by the constant saturation method. Note that the scale of Figures 24 through 31 is not the same for each figure to better illustrate changes in each signal. Figures 30 and 31 illustrate the effect of correlation cancellation using the primary reference signal $s'(t)$ as determined by a reference processor of the present invention utilizing the constant saturation method. Only segments 30b and 31b are dominated by motion induced noise as were segments 24b, and 25b of the measured signals. Additionally, segments 30a, 31a, 30c, and 31c are nearly zero corresponding to the measured signal segments 24a, 25a, 24c, and 25c where there was no motion induced noise.

METHOD FOR ESTIMATING PRIMARY AND SECONDARY

SIGNAL PORTIONS OF

MEASURED SIGNALS IN A PULSE OXIMETER

A copy of a computer subroutine, written in the C programming language, calculates a primary reference $s'(t)$ and a secondary reference $n'(t)$ using the ratiometric method and, using a joint process estimator 60, estimates either the primary or secondary signal portions of two measured signals, each having a primary signal which is

correlated with the primary reference $s'(t)$ and having a secondary signal which is correlated with the secondary reference $n'(t)$, is appended in Appendix A. For example, $S_{\lambda a}(t) = S_{\lambda red}(t) = S_{\lambda 660nm}(t)$ and $S_{\lambda b}(t) = S_{\lambda IR}(t) = S_{\lambda 940nm}(t)$ can be input to the computer subroutine. This subroutine is one way to implement the steps illustrated in the flowchart of Figure 9 for a monitor particularly adapted for pulse oximetry.

The program estimates either the primary signal portions or the secondary signal portions of two light energy signals, one preferably corresponding to light in the visible red range and the other preferably corresponding to light in the infrared range such that a determination of the amount of oxygen, or the saturation of oxygen in the arterial and venous blood components, may be made. The calculation of the saturation is performed in a separate subroutine.

Using the ratiometric method three signals $S_{\lambda a}(t)$, $S_{\lambda b}(t)$ and $S_{\lambda c}(t)$ are input to the subroutine. $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ are used to calculate either the primary or secondary reference signal $s'(t)$ or $n'(t)$. As described above, the wavelengths of light at which $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ are measured are chosen to satisfy the relationship of equation (96). Once either the secondary reference signal $n'(t)$ or the primary reference signal $s'(t)$ is determined, either the primary signal portions $s_{\lambda a}(t)$ and $s_{\lambda c}(t)$ or the secondary signal portions $n_{\lambda a}(t)$ and $n_{\lambda c}(t)$ of the measured signals $S_{\lambda a}(t)$ and $S_{\lambda c}(t)$ are estimated for use in calculation of the oxygen saturation.

The correspondence of the program variables to the variables defined in the discussion of the joint process estimator is as follows:

5 $\Delta_m(t) = nc[m].Delta$
5 $r_{f,m}(t) = nc[m].fref$
5 $r_{b,m}(t) = nc[m].bref$
10 $f_m(t) = nc[m].ferr$
10 $b_m(t) = nc[m].berr$
15 $\Im_m(t) = nc[m].Fswsqr$
15 $\beta_m(t) = nc[m].Bswsqr$
20 $\gamma_m(t) = nc[m].Gamma$
20 $\rho_m, \lambda_a(t) = nc[m].Roh_a$
20 $\rho_m, \lambda_c(t) = nc[m].Roh_c$
25 $e_m, \lambda_a(t) = nc[m].err_a$
25 $e_m, \lambda_c(t) = nc[m].err_c$
25 $\kappa_m, \lambda_a(t) = nc[m].K_a$
30 $\kappa_m, \lambda_c(t) = nc[m].K_c$

30 A first portion of the program performs the initialization of the registers 90, 92, 96, and 98 and intermediate variable values as in the "INITIALIZE CORRELATION CANCELER" box 120 and equations (52) through (56) and equations (73), (74), (77), and (78). A second portion of the program performs the time updates of the delay element variables 110 where the value at the input of each delay element variable 110 is stored in the delay element variable 110 as in the "TIME UPDATE OF $[Z^{-1}]$ ELEMENTS" box 130.

40 A third portion of the program calculates the

reference signal, as in the "CALCULATE SECONDARY REFERENCE (n'(t)) OR PRIMARY REFERENCE (s'(t)) FOR TWO MEASURED SIGNAL SAMPLES" box 140 using the proportionality constant ω_{av} determined by the ratiometric method as in equation (25).

A fourth portion of the program performs the zero-stage update as in the "ZERO-STAGE UPDATE" box 150 where the zero-stage forward prediction error $f_0(t)$ and the zero-stage backward prediction error $b_0(t)$ are set equal to the value of the reference signal $n'(t)$ or $s'(t)$ just calculated. Additionally, zero-stage values of intermediate variables $\beta_0(t)$ and $\beta_0(t)$ ($nc[m].Fwsqr$ and $nc[m].Bwsqr$ in the program) are calculated for use in setting register 90, 92, 96, and 98 values in the least-squares lattice predictor 70 and the regression filters 80a and 80b.

A fifth portion of the program is an iterative loop wherein the loop counter, m , is reset to zero with a maximum of $m = NC_CELLS$, as in the " $m=0$ " box 160 in Figure 9. NC_CELLS is a predetermined maximum value of iterations for the loop. A typical value of NC_CELLS is between 6 and 10, for example. The conditions of the loop are set such that the loop iterates a minimum of five times and continues to iterate until a test for conversion is met or $m = NC_CELLS$. The test for conversion is whether or not the sum of the weighted sum of forward prediction errors plus the weighted sum of backward prediction errors is less than a small number, typically 0.00001 (i.e., $\beta_m(t) + \beta_m(t) \leq 0.00001$).

A sixth portion of the program calculates the forward and backward reflection coefficient $\Gamma_{m,f}(t)$ and $\Gamma_{m,b}(t)$

register 90 and 92 values (nc[m].fref and nc[m].bref in the program) as in the "ORDER UPDATE m^{th} -STAGE OF LSL-PREDICTOR" box 170 and equations (61) and (62). Then forward and backward prediction errors $f_m(t)$ and $b_m(t)$ (nc[m].ferr and nc[m].berr in the program) are calculated as, in equations (63) and (64). Additionally, intermediate variables $\beta_m(t)$, $\beta_m(t)$ and $\gamma_m(t)$ (nc[m].Fswsqr, nc[m].Bswsqr, nc[m].Gamma in the program) are calculated, as in equations (65), (66), and (67). The first cycle of the loop uses the values for nc[0].Fswsqr and nc[0].Bswsqr calculated in the ZERO-STAGE UPDATE portion of the program.

A seventh portion of the program, still within the loop, calculates the regression coefficient $\kappa_m, \lambda_a(t)$ and $\kappa_m, \lambda_c(t)$ register 96 and 98 values (nc[m].K_a and nc[m].K_c in the program) in both regression filters, as in the "ORDER UPDATE m^{th} STAGE OF REGRESSION FILTER(S)" box 180 and equations (68) through (80). Intermediate error signals and variables $e_m, \lambda_a(t)$, $e_m, \lambda_c(t)$, $\rho_m, \lambda_a(t)$, and $\rho_m, \lambda_c(t)$ (nc[m].err_a and nc[m].err_c, nc[m].roh_a, and nc[m].roh_c in the subroutine) are also calculated as in equations (75), (76), (71), and (72), respectively.

The test for convergence of the joint process estimator is performed each time the loop iterates, analogously to the "DONE" box 190. If the sum of the weighted sums of the forward and backward prediction errors $\beta_m(t) + \beta_m(t)$ is less than or equal to 0.00001, the loop terminates. Otherwise, the sixth and seventh portions of the program repeat.

When either the convergence test is passed or $m = \text{NC_CELLS}$, an eighth portion of the program calculates the output of the joint process estimator 60 as in the

"CALCULATE OUTPUT" box 200. This output is a good approximation to both of the primary signals $s''\lambda_a(t)$ and $s''\lambda_c(t)$ or the secondary signals $n''\lambda_a(t)$ and $n''\lambda_c(t)$ for the set of samples $S_{\lambda_a}(t)$ and $S_{\lambda_c}(t)$, input to the program. After many sets of samples are processed by the joint process estimator, a compilation of the outputs provides output waves which are good approximations to the plethysmographic wave or motion artifact at each wavelength, λ_a and λ_c .

Another copy of a computer program subroutine, written in the C programming language, which calculates either a primary reference $s'(t)$ or a secondary reference $n'(t)$ using the constant saturation method and, using a joint process estimator 60, estimates a good approximation to either the primary signal portions or secondary signal portions of two measured signals, each having a primary portion which is correlated to the primary reference signal $s'(t)$ and a secondary portion which is correlated to the secondary reference signal $n'(t)$ and each having been used to calculate the reference signals $s'(t)$ and $n'(t)$, is appended in Appendix B. This subroutine is another way to implement the steps illustrated in the flowchart of Figure 9 for a monitor particularly adapted for pulse oximetry. The two signals are measured at two different wavelengths λ_a and λ_b , where λ_a is typically in the visible region and λ_b is typically in the infrared region. For example, in one embodiment of the present invention, tailored specifically to perform pulse oximetry using the constant saturation method, $\lambda_a = 660\text{nm}$ and $\lambda_b = 940\text{nm}$.

The correspondence of the program variables to the variables defined in the discussion of the joint process estimator is as follows:

```

5       $\Delta_m(t) = nc[m].Delta$ 
       $\Gamma_{f,m}(t) = nc[m].fref$ 
       $\Gamma_{b,m}(t) = nc[m].bref$ 
       $f_m(t) = nc[m].ferr$ 
10      $b_m(t) = nc[m].berr$ 
       $\bar{J}_m(t) = nc[m].Fswsqr$ 
       $\beta_m(t) = nc[m].Bswsqr$ 
15      $\gamma(t) = nc[m].Gamma$ 
       $\rho_{m,\lambda a}(t) = nc[m].Roh_a$ 
20      $\rho_{m,\lambda b}(t) = nc[m].Roh_b$ 
       $e_{m,\lambda a}(t) = nc[m].err_a$ 
       $e_{m,\lambda b}(t) = nc[m].err_b$ 
25      $\kappa_{m,\lambda a}(t) = nc[m].K_a$ 
       $\kappa_{m,\lambda b}(t) = nc[m].K_b$ 

```

30 First and second portions of the subroutine are the
35 same as the first and second portions of the above
40 described subroutine tailored for the ratiometric method of
45 determining either the primary reference $s'(t)$ or the noise
50 reference $n'(t)$. The calculation of saturation is performed
55 in a separate module. Various methods for calculation of
60 the oxygen saturation are known to those skilled in the
65 art. One such calculation is described in the articles by
70 G.A. Mook, et al, and Michael R. Neuman cited above. Once
75 the concentration of oxygenated hemoglobin and deoxygenated
80 hemoglobin are determined, the value of the saturation is
85 determined similarly to equations (85) through (92) wherein

measurements at times t_1 and t_2 are made at different, yet proximate times over which the saturation is relatively constant. For pulse oximetry, the average saturation at time $t = (t_1+t_2)/2$ is then determined by:

$$5 \quad \text{Saturation}_{\text{Art}}(t) = c_{\text{HbO}_2}^A(t) / [c_{\text{HbO}_2}^A(t) + c_{\text{Hb}}^A(t)] \quad (107a)$$

$$10 \quad = \frac{\epsilon_{\text{Hb}, \lambda_a} - \epsilon_{\text{Hb}, \lambda_b} (\Delta s_{\lambda_a} / \Delta s_{\lambda_b})}{\epsilon_{\text{Hb}, \lambda_a} - \epsilon_{\text{HbO}_2, \lambda_a} - (\epsilon_{\text{Hb}, \lambda_b} - \epsilon_{\text{HbO}_2, \lambda_b}) (\Delta s_{\lambda_a} / \Delta s_{\lambda_b})} \quad (107b)$$

$$15 \quad \text{Saturation}_{\text{Ven}}(t) = c_{\text{HbO}_2}^V(t) / [c_{\text{HbO}_2}^V(t) + c_{\text{HB}}^V(t)] \quad (108a)$$

$$= \frac{\epsilon_{\text{Hb}, \lambda_a} - \epsilon_{\text{Hb}, \lambda_b} (\Delta n_{\lambda_a} / \Delta n_{\lambda_b})}{\epsilon_{\text{Hb}, \lambda_a} - \epsilon_{\text{HbO}_2, \lambda_a} - (\epsilon_{\text{Hb}, \lambda_b} - \epsilon_{\text{HbO}_2, \lambda_b}) (\Delta n_{\lambda_a} / \Delta n_{\lambda_b})} \quad (108b)$$

20 A third portion of the subroutine calculates either the primary reference or secondary reference, as in the "CALCULATE PRIMARY OR SECONDARY REFERENCE ($s'(t)$ or $n'(t)$) FOR TWO MEASURED SIGNAL SAMPLES" box 140 for the signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ using the proportionality constants $\omega_a(t)$ and $\omega_v(t)$ determined by the constant saturation method as in equation (3). The saturation is calculated in a separate subroutine and a value of $\omega_a(t)$ or $\omega_v(t)$ is imported to the present subroutine for estimating either the primary portions $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ or the secondary portions $n_{\lambda_a}(t)$ and $n_{\lambda_b}(t)$ of the composite measured signals $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$.

25 Fourth, fifth, and sixth portions of the subroutine are similar to the fourth, fifth, and sixth portions of the above described program tailored for the ratiometric method. However, the signals being used to estimate the primary signal portions $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ or the secondary signal portions $n_{\lambda_a}(t)$ and $n_{\lambda_b}(t)$ in the present subroutine

tailored for the constant saturation method, are $S_{\lambda_a}(t)$ and $S_{\lambda_b}(t)$, the same signals that were used to calculate the reference signal $s'(t)$ or $n'(t)$.

A seventh portion of the program, still within the loop begun in the fifth portion of the program, calculates the regression coefficient register 96 and 98 values $\kappa_m, \lambda_a(t)$ and $\kappa_m, \lambda_b(t)$ ($nc[m].K_a$ and $nc[m].K_b$ in the program) in both regression filters, as in the "ORDER UPDATE mth STAGE OF REGRESSION FILTER(S)" box 180 and equations (68) through (80). Intermediate error signals and variables $e_m, \lambda_a(t)$, $e_m, \lambda_b(t)$, $\rho_m, \lambda_a(t)$, and $\rho_m, \lambda_b(t)$ ($nc[m].err_a$ and $nc[m].err_b$, $nc[m].roh_a$, and $nc[m].roh_b$ in the subroutine) are also calculated as in equations (70), (75), (68), and (71), respectively.

The loop iterates until the test for convergence is passed, the test being the same as described above for the subroutine tailored for the ratiometric method. The output of the present subroutine is a good approximation to the primary signals $s''\lambda_a(t)$ and $s''\lambda_b(t)$ or the secondary signals $n''\lambda_a(t)$ and $n''\lambda_b(t)$ for the set of samples $S_{\lambda_a}(t)$ and $S_{\lambda_b}(t)$ input to the program. After approximations to the primary signal portions or the secondary signals portions of many sets of measured signal samples are estimated by the joint process estimator, a compilation of the outputs provides waves which are good approximations to the plethysmographic wave or motion artifact at each wavelength, λ_a and λ_b . The estimating process of the iterative loop is the same in either subroutine, only the sample values $S_{\lambda_a}(t)$ and $S_{\lambda_c}(t)$ or $S_{\lambda_a}(t)$ and $S_{\lambda_b}(t)$ input to the subroutine for use in estimation of the primary signal portions $s_{\lambda_a}(t)$ and $s_{\lambda_c}(t)$ or $s_{\lambda_a}(t)$ and $s_{\lambda_b}(t)$ or of the secondary signal portions $n_{\lambda_a}(t)$ and $n_{\lambda_c}(t)$ or

$n\lambda_a(t)$ and $n\lambda_b(t)$ and how the primary and secondary reference signals $s'(t)$ and $n'(t)$ are calculated are different for the ratiometric method and the constant saturation methods.

Independent of the method used, ratiometric or constant saturation, the approximations to either the primary signal values or the secondary signal values are input to a separate subroutine in which the saturation of oxygen in the arterial and venous blood is calculated. If the constant saturation method is used, the saturation calculation subroutine also determines values for the proportionality constants $\omega_a(t)$ and $\omega_v(t)$ as defined in equation (3) and discussed above. The concentration of oxygenated arterial and venous blood can be found from the approximations to the primary or secondary signal values since they are made up of terms comprising $x(t)$, the thickness of arterial and venous blood in the finger; absorption coefficients of oxygenated and de-oxygenated hemoglobin, at each measured wavelength; and $c_{HbO_2}(t)$ and $c_{Hb}(t)$, the concentrations of oxygenated and de-oxygenated hemoglobin, respectively. The saturation is a ratio of the concentration of one constituent, A_5 , with respect to the total concentration of constituents in the volume containing A_5 and A_6 or the ratio of the concentration of one constituent A_3 , with respect to the total concentration of constituents in the volume containing A_3 and A_4 . Thus, the thickness, $x(t)$, is divided out of the saturation calculation and need not be predetermined. Additionally, the absorption coefficients are constant at each wavelength. The saturation of oxygenated arterial and venous blood is then determined as in equations (107) and (108).

5 While one embodiment of a physiological monitor incorporating a processor of the present invention for determining a reference signal for use in a correlation canceler, such as an adaptive noise canceler, to remove or derive primary and secondary components from a physiological measurement has been described in the form of a pulse oximeter, it will be obvious to one skilled in the art that other types of physiological monitors may also employ the above described techniques.

10 Furthermore, the signal processing techniques described in the present invention may be used to compute the arterial and venous blood oxygen saturations of a physiological system on a continuous or nearly continuous time basis. These calculations may be performed, regardless of whether or not the physiological system undergoes voluntary motion. The arterial pulsation induced primary plethysmographic signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ may be used to compute arterial blood oxygen saturation. The primary signals $s_{\lambda a}(t)$ and $s_{\lambda b}(t)$ can always be introduced into the measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ if at least two requirements are met. The two requirements include the selection of two or more flesh penetrating and blood absorbing wavelengths which are optically modulated by the arterial pulsation and an instrument design which passes all or portions of all electromagnetic signals which are related to the pulsation. Similarly, the secondary signals $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ related to venous blood flow may be used to compute its corresponding oxygen saturation. The secondary signal components $n_{\lambda a}(t)$ and $n_{\lambda b}(t)$ can be guaranteed to be contained in the measured signals $S_{\lambda a}(t)$ and $S_{\lambda b}(t)$ if the two or more flesh penetrating and blood absorbing wavelengths are processed to pass all or portions of all electromagnetic signals relating to venous blood

flow. This may include but is not limited to all or portions of all signals which are related to the involuntary action of breathing. Similarly, it must be understood that there are many different types of physical systems which may be configured to yield two or more measurement signals each possessing a primary and secondary signal portion. In a great many of such physical systems it will be possible to derive one or more reference signals. The reference signals may be used in conjunction with a correlation canceler, such as an adaptive noise canceler, to derive either the primary and/or secondary signal components of the two or more measurement signals on a continuous or intermittent time basis.

Another embodiment of a physiological monitor incorporating a processor of the present invention for determining a reference signal for use in a correlation canceler, such as an adaptive noise canceler, to remove or derive primary and secondary components from a physiological measurement may be described in the form of a instrument which measures blood pressure. There are several ways of obtaining blood pressure measurements, such as tonometry, and pulse wave velocity. Both of these methods are substantially related to plethysmography.

Tonometry is a measurement method in which a direct reading of the arterial pressure pulse is made non-invasively. These measurements are invariably made through the use of a piezoelectric force transducer, the surface of which is gently pressed against a near-surface artery supported by underlying bone. If the transducer is sufficiently pressed against the artery that its surface is in complete contact with the tissue; then, knowing its surface area, its output can be directly read as pressure. This "flattening" of the

arterial wall leads to the name of this method, applanation tonometry. The pulse wave velocity technique relies on the concept that the speed with which the pressure pulse, generated at the heart, travels "down" the arterial system is dependent on pressure. In each of these cases plethysmographic waveforms are used to determine the blood pressure of a patient.

Furthermore, it will be understood that transformations of measured signals other than logarithmic conversion and determination of a proportionality factor which allows removal or derivation of the primary or secondary signal portions for determination of a reference signal are possible. Additionally, although the proportionality factor ω has been described herein as a ratio of a portion of a first signal to a portion of a second signal, a similar proportionality constant determined as a ratio of a portion of a second signal to a portion of a first signal could equally well be utilized in the processor of the present invention. In the latter case, a secondary reference signal would generally resemble $n'(t) = n\lambda_b(t) - \omega n\lambda_a(t)$.

Furthermore, it will be understood that correlation cancellation techniques other than joint process estimation may be used together with the reference signals of the present invention. These may include but are not limited to least mean square algorithms, wavelet transforms, spectral estimation techniques, neural networks, Weiner filters, Kalman filters, QR-decomposition based algorithms among others. The implementation that we feel is the best, as of this filing, is the normalized least square lattice algorithm an implementation of which is listed in Appendix C.

5 It will also be obvious to one skilled in the art that
for most physiological measurements, two wavelengths may be
determined which will enable a signal to be measured which
is indicative of a quantity of a component about which
information is desired. Information about a constituent of
any energy absorbing physiological material may be
determined by a physiological monitor incorporating a
signal processor of the present invention and an
correlation canceler by determining wavelengths which are
absorbed primarily by the constituent of interest. For most
physiological measurements, this is a simple determination.

10
15 Moreover, one skilled in the art will realize that any
portion of a patient or a material derived from a patient
may be used to take measurements for a physiological
monitor incorporating a processor of the present invention
and a correlation canceler. Such areas include a digit such
as a finger, but are not limited to a finger.

20
25 One skilled in the art will realize that many
different types of physiological monitors may employ a
signal processor of the present invention in conjunction
with a correlation canceler, such as an adaptive noise
canceler. Other types of physiological monitors include,
but are in not limited to, electron cardiographs, blood
pressure monitors, blood gas saturation (other than oxygen
saturation) monitors, capnographs, heart rate monitors,
respiration monitors, or depth of anesthesia monitors.
30 Additionally, monitors which measure the pressure and
quantity of a substance within the body such as a
breathalyzer, a drug monitor, a cholesterol monitor, a
glucose monitor, a carbon dioxide monitor, a glucose
monitor, or a carbon monoxide monitor may also employ the

above described techniques for removal of primary or secondary signal portions.

Furthermore, one skilled in the art will realize that the above described techniques of primary or secondary signal removal or derivation from a composite signal including both primary and secondary components can also be performed on electrocardiography (ECG) signals which are derived from positions on the body which are close and highly correlated to each other. It must be understood that a tripolar Laplacian electrode sensor such as that depicted in Figure 32 which is a modification of a bipolar Laplacian electrode sensor discussed in the article "Body Surface Laplacian ECG Mapping" by Bin He and Richard J. Cohen contained in the journal IEEE Transactions on Biomedical Engineering, Vol. 39, No. 11, November 1992 could be used as an ECG sensor. This article is hereby incorporated as reference. It must also be understood that there are a myriad of possible ECG sensor geometry's that may be used to satisfy the requirements of the present invention.

Furthermore, one skilled in the art will realize that the above described techniques of primary or secondary signal removal or derivation from a composite signal including both primary and secondary components can also be performed on signals made up of reflected energy, rather than transmitted energy. One skilled in the art will also realize that a primary or secondary portion of a measured signal of any type of energy, including but not limited to sound energy, X-ray energy, gamma ray energy, or light energy can be estimated by the techniques described above. Thus, one skilled in the art will realize that the processor of the present invention and a correlation canceler can be applied in such monitors as those using

5 ultrasound where a signal is transmitted through a portion of the body and reflected back from within the body back through this portion of the body. Additionally, monitors such as echo cardiographs may also utilize the techniques
10 of the present invention since they too rely on transmission and reflection.

10
15 While the present invention has been described in terms of a physiological monitor, one skilled in the art will realize that the signal processing techniques of the present invention can be applied in many areas, including but not limited to the processing of a physiological signal. The present invention may be applied in any situation where a signal processor comprising a detector receives a first signal which includes a first primary signal portion and a first secondary signal portion and a second signal which includes a second primary signal portion and a second secondary signal portion. The first and second signals propagate through a common medium and the first and second primary signal portions are correlated with one another. Additionally, at least a portion of the first and second secondary signal portions are correlated with one another due to a perturbation of the medium while the first and second signals are propagating through the medium. The processor receives the first and second signals and may combine the first and second signals to generate a secondary reference in which is uncorrelated with the primary signal portions of the measured signals or a primary reference which is uncorrelated with the secondary signal portions of the measured signals. Thus, the signal processor of the present invention is readily applicable to numerous signal processing areas.
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25
30

CLAIMS

1. In combination:

detector means for receiving a first signal which travels along a first propagation path and a second signal which travels along a second propagation path, a portion of said first and second propagation paths being located in the same propagation medium, wherein said first signal has a primary signal portion and a secondary signal portion and said second signal has a primary signal portion and a secondary signal portion; and

10 first signal processor means having input means for receiving said first and second signals and combining said first and second signals for generating either a primary or secondary reference signal having a significant component which is a function of either of, respectively, said primary or said secondary signal portions of said first and second signals.

2. The combination recited in claim 1, further comprising second signal processor means for receiving said secondary reference signal and said first signal and for deriving therefrom an output signal having a significant component which is a function of said primary signal portion of said first signal.

3. The combination recited in claim 2, wherein said second signal processor means includes a correlation canceler.

4. The combination recited in claim 2, wherein said second signal processor means includes an adaptive noise canceler.

5. The combination recited in claim 4, wherein said adaptive noise canceler includes a joint process estimator.

6. The combination recited in claim 5, wherein said joint process estimator comprises a least-squares lattice predictor and a regression filter.

7. The combination recited in claim 1, further comprising second signal processor means for receiving said primary reference signal and said first signal and for deriving therefrom an output signal having a significant component which is a function of said secondary signal portion of said first signal.

8. The combination recited in claim 1, further comprising second signal processor means for receiving said secondary reference signal and said first signal and for deriving therefrom an output signal having a significant component which is a function of said primary signal portion of said second signal.

9. The combination recited in claim 1, further comprising second signal processor means for receiving said primary reference signal and said first signal and for deriving therefrom an output signal having a significant component which is a function of said secondary signal portion of said second signal.

10. The combination recited in claim 1, wherein said detector means includes a sensor for sensing a physiological function depending upon said first and second signals.

11. The combination recited in claim 10, wherein said sensor is adapted to measure a blood constituent.

12. The combination recited in claim 11, wherein the blood constituent measured by said sensor is blood gas.

13. The combination recited in claim 10, wherein said sensor includes a device that is responsive to electromagnetic energy.

14. The combination recited in claim 1, further comprising electromagnetic means connected to said detector means for measuring a plethysmographic waveform depending upon said first and second signals received by said detector means through said propagation medium, said propagation medium including living tissue.

15. The combination recited in claim 1, further comprising pulse oximeter means connected to said detector means for measuring a physiological condition depending upon said first and second signals received by said detector means through said propagation medium, said propagation medium including living tissue.

16. The combination recited in claim 1, further comprising blood pressure monitoring means connected to said detector means for measuring a physiological condition depending upon said first and second signals received by said detector means through said propagation medium, said propagation medium including living tissue.

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17. The combination recited in claim 1, further comprising electrocardiogram means connected to said detector means for measuring a physiological condition depending upon said first and second signals received by said detector means through said propagation medium, said propagation medium including living tissue.

18. The combination recited in claim 17, wherein said electrocardiogram means for measuring a physiological condition includes a tripolar electrode sensor having three concentrically arranged electrodes.

19. Apparatus for indicating the constituency of a material, said apparatus comprising:

first signal processor means for receiving first and second inputs, the first input comprising one of a plurality of reference signals, each of said plurality of reference signals being related to the second input;

means for applying said second input and each of said plurality of reference signals to said first signal processor means; and

means for detecting the output signals of said first signal processor means for each of said plurality of reference signals applied to said signal processor means, wherein said output signals are indicative of the constituency of said material.

20. The apparatus recited in claim 19, wherein said first signal processor means is a correlation canceler.

21. The apparatus recited in claim 19, wherein said first signal processor means is an adaptive noise canceler.

22. The apparatus recited in claim 21, wherein said adaptive noise canceler comprises a joint process estimator.

23. The apparatus recited in claim 22, wherein said joint process estimator comprises a least-squares lattice predictor and a regression filter.

24. The apparatus recited in claim 19, further comprising second signal processor means receiving each of the output signals of said first signal processor means, said second signal processor means integrating each of said output signals for forming a cumulative output signal which is indicative of said integrated output signals.

5

25. The apparatus recited in claim 24, wherein said detecting means is responsive to the extremum of said cumulative output signal of said second signal processor means for providing an indication of the constituency of the material.

5

26. The apparatus recited in claim 24, wherein said detecting means is responsive to the inflections of said cumulative output signal of said second signal processor means for providing an indication of the constituency of the material.

27. The apparatus recited in claim 19, wherein said second input applied to said first signal processor means is one of a first or a second signal, each of said first and second signals having an arterial signal portion and another signal portion that is indicative of venous blood, and further comprising second signal processor means for receiving said first and second signals and a plurality of signal coefficients, wherein the output of said second signal processor means forms said first input applied to said first signal processor means, the output of said second signal processor means including a first component which is related to the arterial signal portions of said first and second signals and a second component which is related to the other signal portions of said first and second signals.

28. The apparatus recited in claim 27, wherein the other signal portion of each of said first and second signals includes an indication of human respiration.

29. The apparatus recited in claim 27, wherein at least one of said plurality of signal coefficients received by said second signal processor means relates to the arterial signal portions of said first and second signals of said second input to said first signal processor.

30. The apparatus recited in claim 27, wherein at least one of said plurality of signal coefficients received by said second signal processor means relates to the other signal portions of said first and second signals of said second input to said first signal processor means.

31. The apparatus recited in claim 19, wherein said second input applied to said first signal processor means is one of a first or a second signal, each of said first and second signals having an arterial signal portion and another signal portion that is indicative of venous blood, and further comprising second signal processor means for receiving said first and second signals and a plurality of signal coefficients, wherein the output of said second signal processor means forms said first input applied to said first signal processor means, the output of said second signal processor means including a component which is related to the arterial signal portions of said first and second signals or a component which is related to the other signal portions of said first and second signals.

32. The apparatus recited in claim 31, wherein the other signal portion of each of said first and second signals includes an indication of human respiration.

33. The apparatus recited in claim 19, wherein said material is human tissue.

34. The apparatus recited in claim 19, wherein said detecting means is responsive to the power of the output signals of said first signal processor means for indicating the constituency of said material.

35. Apparatus for computing arterial and venous signals in living tissue, said apparatus comprising:

5 detector means for receiving a first signal which travels along a first propagation path and a second signal which travels along a second propagation path, a portion of said first and second propagation paths being located in a propagation medium, wherein said first signal has an arterial signal portion that is indicative of arterial blood and another signal portion that is indicative of venous blood, and said second signal has an arterial signal portion that is indicative of arterial blood and another signal portion that is indicative of venous blood; and

10 signal processor means having input means for receiving said first and second signals and combining said first and

second signals to generate a signal having a significant component which is a function of either of said arterial or said other signal portions of said first and second signal.

36. The apparatus recited in claim 35, wherein the other signal portion of each of said first and second signals includes an indication of human respiration.

37. Apparatus for computing arterial and venous blood constituent values in living tissue, said apparatus comprising:

signal processor means for receiving first and second inputs, a first of said inputs comprising one of a plurality of reference signals, each of said plurality of reference signals being related to the second input;

means for applying said second input and each of said plurality of reference signals of said first input to said signal processor means; and

means for detecting the power of the output signals of said signal processor means for each of said plurality of reference signals of said first input, wherein said power is indicative of said arterial and venous blood constituent values.

38. The apparatus recited in claim 37, wherein said arterial and venous blood constituent values are the oxygen saturation of arterial and venous blood, respectively.

SIGNAL PROCESSING APPARATUS

Abstract of the Invention

A signal processor which acquires a first signal, including a first primary signal portion and a first secondary signal portion, and a second signal, including a second primary signal portion and a second secondary signal portion, wherein the first and second primary signal portions are correlated. The signals may be acquired by propagating energy through a medium and measuring an attenuated signal after transmission or reflection. Alternatively, the signals may be acquired by measuring energy generated by the medium. A processor of the present invention generates a primary or secondary reference signal which is a combination, respectively, of only the primary or secondary signal portions. The secondary reference signal is then used to remove the secondary portion of each of the first and second measured signals via a correlation canceler, such as an adaptive noise canceler, preferably of the joint process estimator type. The primary reference signal is used to remove the primary portion of each of the first and second measured signals via a correlation canceler. The processor of the present invention may be employed in conjunction with a correlation canceler in physiological monitors wherein the known properties of energy attenuation through a medium are used to determine physiological characteristics of the medium. Many physiological conditions, such as the pulse, or blood pressure of a patient or the concentration of a constituent in a medium, can be determined from the primary or secondary portions of the signal after other signal portion is removed.

```

***** APPENDIX A *****
***** Least Square Lattice *****
***** Noise Cancelling *****
/* Example for ratiometric approach to noise cancelling */
#define LAMBDA 0.95

void OxiLSL_NC( int      reset,
                int      passes,
                int      *signal_1,
                int      *signal_2,
                int      *signal_3,
                int      *target_1,
                int      *target_2) {

    int      i, ii, k, m, n, contraction;
    static int      *s_a, *s_b, *s_c, *out_a, *out_c;
    static float   Delta_sqr, scale, noise_ref;

    if( reset == TRUE){
        s_a      = signal_1;
        s_b      = signal_2;
        s_c      = signal_3;
        out_a   = target_1;
        out_c   = target_2;
        factor  = 1.5;
        scale   = 1.0 /4160.0;
    }
    /* noise canceller initialization at time t=0 */
    nc[0].berr  = 0.0;
    nc[0].Gamma = 1.0;

    for(m=0; m<NC_CELLS; m++) {
        nc[m].err_a  = 0.0;
        nc[m].err_b  = 0.0;
        nc[m].Roh_a  = 0.0;
        nc[m].Roh_c  = 0.0;
        nc[m].Delta  = 0.0;
        nc[m].Fswsqr = 0.00001;
        nc[m].Bswsqr = 0.00001;
    }
}

===== END INITIALIZATION =====

for(k=0; k<passes; k++){
    contraction = FALSE;
    for(m=0; m< NC_CELLS; m++) {          /* Update delay elements */
        nc[m].berri  = nc[m].berr;
        nc[m].Bswsqr1 = nc[m].Bswsqr;
    }

    noise_ref  = factor * log(1.0 - (*s_a) * scale)
    - log(1.0 - (*s_b) * scale);
    nc[0].err_a = log(1.0 - (*s_b) * scale);
    nc[0].err_b = log(1.0 - (*s_c) * scale);
}

```

```

++s_a;
++s_b;
++s_c;

nc[0].ferr = noise_ref ;
nc[0].berr = noise_ref ;
nc[0].Fswsqr = LAMBDA * nc[0].Fswsqr + noise_ref * noise_ref;
nc[0].Bswsqr = nc[0].Fswsqr;

/* Order Update */
for(n=1; ( n < NC_CELLS) && (contraction == FALSE); n++) {

    /* Adaptive Lattice Section */

    m = n-1;
    ii= n-1;

    nc[m].Delta *= LAMBDA;
    nc[m].Delta += nc[m].berri * nc[m].ferr / nc[m].Gamma ;
    Delta_sqr = nc[m].Delta * nc[m].Delta;

    nc[n].fref = -nc[m].Delta / nc[m].Bswsqr1;
    nc[n].bref = -nc[m].Delta / nc[m].Fswsqr;

    nc[n].ferr = nc[m].ferr + nc[n].fref * nc[m].berri;
    nc[n].berr = nc[m].berri + nc[n].bref * nc[m].ferr;

    nc[n].Fswsqr = nc[m].Fswsqr - Delta_sqr / nc[m].Bswsqr1;
    nc[n].Bswsqr = nc[m].Bswsqr1 - Delta_sqr / nc[m].Fswsqr;

    if( (nc[n].Fswsqr + nc[n].Bswsqr) > 0.00001 || (n < 5) ) {
        nc[n].Gamma = nc[m].Gamma - nc[m].berri * nc[m].berri / nc[m].Bswsqr1;
        if(nc[n].Gamma < 0.05) nc[n].Gamma = 0.05;
        if(nc[n].Gamma > 1.00) nc[n].Gamma = 1.00;
    }

    /* Joint Process Estimation Section */

    nc[m].Roh_a *= LAMBDA;
    nc[m].Roh_a += nc[m].berr * nc[m].err_a / nc[m].Gamma ;
    nc[m].k_a = nc[m].Roh_a / nc[m].Bswsqr;
    nc[n].err_a = nc[m].err_a - nc[m].k_a * nc[m].berr;

    nc[m].Roh_c *= LAMBDA;
    nc[m].Roh_c += nc[m].berr * nc[m].err_b / nc[m].Gamma ;
    nc[m].k_c = nc[m].Roh_c / nc[m].Bswsqr;
    nc[n].err_b = nc[m].err_b - nc[m].k_c * nc[m].berr;

}

else {
    contraction = TRUE;
    for(i=n; i<NC_CELLS; i++) {
        nc[i].err_a = 0.0;
        nc[i].Roh_a = 0.0;
        nc[i].err_b = 0.0;
        nc[i].Roh_c = 0.0;
        nc[i].Delta = 0.0;
        nc[i].Fswsqr = 0.00001;
        nc[i].Bswsqr = 0.00001;
        nc[i].Bswsqr1 = 0.00001;
    }
}

```

```
    }
}

{
    *out_a++ = (int)( (-exp(nc[ii].err_a) +1.0) / scale) ;
    *out_c++ = (int)( (-exp(nc[ii].err_b) +1.0) / scale) ;

}
}
***** Least Square Lattice *****
***** ***** ***** ***** *****
***** ***** ***** ***** *****/
```

```

***** APPENDIX B *****
***** Least Square Lattice *****
***** Noise Cancelling *****
/* Example for constant saturation approach to noise cancelling */
#define LAMBDA 0.95

void OxILSL_NC( int      reset,
                int      passes,
                int      sat_factor,
                int      *signal_1,
                int      *signal_2,
                int      *target_1,
                int      *target_2) {

    int      i, ii, k, m, n, contraction;
static int      *s_a, *s_b, *out_a, *out_b;
static float     Delta_sqr, scale, noise_ref;

if( reset == TRUE){
    s_a = signal_1;
    s_b = signal_2;
    out_a = target_1;
    out_b = target_2;
    scale = 1.0 /4160.0;

/* noise canceller initialization at time t=0 */

    nc[0].berr = 0.0;
    nc[0].Gamma = 1.0;

    for(m=0; m<NC_CELLS; m++) {
        nc[m].err_a = 0.0;
        nc[m].err_b = 0.0;
        nc[m].Roh_a = 0.0;
        nc[m].Roh_b = 0.0;
        nc[m].Delta = 0.0;
        nc[m].Fwsqr = 0.00001;
        nc[m].Bwsqr = 0.00001;
    }
}

/*----- END INITIALIZATION -----*/
for(k=0; k<passes; k++){

    contraction = FALSE;
    for(m=0; m< NC_CELLS; m++) { /* Update delay elements */
        nc[m].berri = nc[m].berr;
        nc[m].Bwsqr1 = nc[m].Bwsqr;
    }

    noise_ref = sat_factor * log(1.0 - (*s_a) * scale)
               - log(1.0 - (*s_b) * scale);
    nc[0].err_a = log(1.0 - (*s_a) * scale);
    nc[0].err_b = log(1.0 - (*s_b) * scale);

    ++s_a;
}

```

```

++s_b;

nc[0].ferr = noise_ref ;
nc[0].berr = noise_ref ;
nc[0].Fwsqr = LAMBDA * nc[0].Fwsqr + noise_ref * noise_ref;
nc[0].Bwsqr = nc[0].Fwsqr;

/* Order Update */
for(n=1; ( n < NC_CELLS) && (contraction == FALSE); n++) {

    /* Adaptive Lattice Section */

    m = n-1;
    ii= n-1;

    nc[m].Delta *= LAMBDA;
    nc[m].Delta += nc[m].berri * nc[m].ferr / nc[m].Gamma ;
    Delta_sqr = nc[m].Delta * nc[m].Delta;

    nc[n].fref = -nc[m].Delta / nc[m].Bwsqr1;
    nc[n].bref = -nc[m].Delta / nc[m].Fwsqr;

    nc[n].ferr = nc[m].ferr + nc[n].fref * nc[m].berri;
    nc[n].berr = nc[m].berri + nc[n].bref * nc[m].ferr;

    nc[n].Fwsqr = nc[m].Fwsqr - Delta_sqr / nc[m].Bwsqr1;
    nc[n].Bwsqr = nc[m].Bwsqr1 - Delta_sqr / nc[m].Fwsqr;

    if( (nc[n].Fwsqr + nc[n].Bwsqr) > 0.00001 || (n < 5) ) {
        nc[n].Gamma = nc[m].Gamma - nc[m].berri * nc[m].berri / nc[m].Bwsqr1;
        if(nc[n].Gamma < 0.05) nc[n].Gamma = 0.05;
        if(nc[n].Gamma > 1.00) nc[n].Gamma = 1.00;
    }

    /* Joint Process Estimation Section */

    nc[m].Roh_a *= LAMBDA;
    nc[m].Roh_a += nc[m].berr * nc[m].err_a / nc[m].Gamma ;
    nc[m].k_a = nc[m].Roh_a / nc[m].Bwsqr;
    nc[n].err_a = nc[m].err_a - nc[m].k_a * nc[m].berr;

    nc[m].Roh_b *= LAMBDA;
    nc[m].Roh_b += nc[m].berr * nc[m].err_b / nc[m].Gamma ;
    nc[m].k_b = nc[m].Roh_b / nc[m].Bwsqr;
    nc[n].err_b = nc[m].err_b - nc[m].k_b * nc[m].berr;

}

else {
    contraction = TRUE;
    for(i=n; i<NC_CELLS; i++) {
        nc[i].err_a = 0.0;
        nc[i].Roh_a = 0.0;
        nc[i].err_b = 0.0;
        nc[i].Roh_b = 0.0;
        nc[i].Delta = 0.0;
        nc[i].Fwsqr = 0.00001;
        nc[i].Bwsqr = 0.00001;
        nc[i].Bwsqr1 = 0.00001;
    }
}

```

```
    }
}

*out_a++ = (int)( (-exp(nc[ii].err_a) +1.0) / scale) ;
*out_c++ = (int)( (-exp(nc[ii].err_b) +1.0) / scale) ;

}

***** Least Square Lattice *****
```

/* -----
Copyright (c) Masimo Corporation (tm) 1992, 1993. All Rights Reserved.

File: smanc.c1

Description: Improved Normalized Least Squares Lattice ANC

Public Functions: SANC_Calc
SANC_Init

Notes:

This version uses many of the same optimization techniques as the .asm version.

History:

HGK 04/29/93 Design Note SDN43 Rev A

*/

```
#define MODULE_ID 1007
```

```
#include <masimo.h> /* platform descriptions */
```

```
#include <math.h>
```

```
#include <smanc.h> /* self */
```

```
#define MAX(a,b) (a) > (b) ? (a) : (b)
```

```
#define MIN(a,b) (a) < (b) ? (a) : (b)
```

```
#define MIN_VAL 0.01
```

```
#define MAX_DEL 0.999999999999999
```

```
#define MIN_DEL -0.999999999999999
```

```
#define MAX_RHO 2.0
```

```
#define MIN_RHO -2.0
```

```
#define MAX_BSERR 1.0
```

```
#define MIN_BSERR 1E-15
```

/* The following macros provide efficient access to the lattice */

```
#define XBERR 0
```

```
#define XBERR_1 1
```

```
#define XDELTA 2
```

```
#define XDELTA_1 3
```

```
#define XGAMMA 4
```

```
#define XGAMMA_1 5
```

```
#define XBSERR 6
```

```
#define XBSERR_1 7
```

```
#define XERR 8
```

```
#define XFERR 9
```

```
#define Xrho 10
```

```
#define berr (*p + XBERR))
```

```
#define P_berr_1 (*p + XBERR_1 - SANC_CELL_SIZE))
```

```
#define P_berr (*p + XBERR - SANC_CELL_SIZE))
```

```
#define berr_1 (*p + XBERR_1))
```

```
#define Bserr (*p + XBSERR))
```

```
#define Bserr_1 (*p + XBSERR_1))
```

```
#define P_Bserr_1 (*p + XBSERR_1 - SANC_CELL_SIZE))
```

```
#define P_delta (*p + XDELTA - SANC_CELL_SIZE))
```

```

#define delta          (* (p + xDELTA))
#define delta_1        (* (p + xDELTA_1))
#define P_delta_1     (* (p + xDELTA_1 - SANC_CELL_SIZE))

#define err            (* (p + xERR))
#define N_err          (* (p + xERR + SANC_CELL_SIZE))

#define P_ferr         (* (p + xFERR - SANC_CELL_SIZE))
#define ferr           (* (p + xFERR))

#define gamma          (* (p + xGAMMA))
#define P_gamma         (* (p + xGAMMA - SANC_CELL_SIZE))
#define N_gamma         (* (p + xGAMMA + SANC_CELL_SIZE))
#define P_gamma_1       (* (p + xGAMMA_1 - SANC_CELL_SIZE))
#define gamma_1         (* (p + xGAMMA_1))

#define rho             (* (p + xRho))

FLOAT32
SANC_Calc(
    SANC_DATA *anc,      /* input, context handle */ 
    FLOAT32 nps,         /* input, noise plus signal */ 
    FLOAT32 noise)       /* input, noise reference */ 
{
    INT32 m;
    FLOAT32 *p;
    FLOAT32 B,F,B2,F2;
    FLOAT32 qd2,qd3;
    INT32 output_cell;
    BOOL Bflag;

    BUG1(anc); BUG1(nps); BUG1(noise);

    /* Update time delay elements in cell structure ----- */
    p = (FLOAT32 *)anc->cells;
    for (m = 0; m <= anc->cc; m++) {
        gamma_1 = gamma;
        berr_1 = berr;
        Bserr_1 = Bserr;
        delta_1 = delta;
        p += SANC_CELL_SIZE;
    }

    /* Handle Cell # 0 ----- */
    p = (FLOAT32 *)anc->cells;
    Bserr = anc->lambda * Bserr_1 + noise * noise;
    Bserr = MAX(Bserr, MIN_BSERR);

    ferr = noise / SQRTF(Bserr);
    ferr = MAX(ferr, MIN_DEL);
    ferr = MIN(ferr, MAX_DEL);

    berr = ferr;

    rho = anc->lambda * SQRTF(Bserr_1 / Bserr) * rho + berr * nps;
    N_err = nps - rho * berr;
}

```

```

/* Initialize Cell voter ----- */
output_cell = anc->cc - 1;      /* Assume last cell for starter */
Bflag = FALSE;

for (m = 1; m < anc->cc; m++) {
    p += SANC_CELL_SIZE;

    B = SQRTF(1.0 - P_berr_1 * P_berr_1);      B2 = 1.0/B;
    F = SQRTF(1.0 - P_ferr_1 * P_ferr_1);      F2 = 1.0/F;

    P_delta = P_delta_1 * F * B + P_berr_1 * P_ferr;
    P_delta = MAX(P_delta, MIN_DEL);
    P_delta = MIN(P_delta, MAX_DEL);
    qd3 = 1.0 - P_delta * P_delta;
    qd2 = 1.0 / SQRTF(qd3);

    ferr = (P_ferr_1 - P_delta * P_berr_1) * qd2 * B2;
    ferr = MAX(ferr, MIN_DEL);
    ferr = MIN(ferr, MAX_DEL);

    berr = (P_berr_1 - P_delta * P_ferr_1) * qd2 * F2;
    berr = MAX(berr, MIN_DEL);
    berr = MIN(berr, MAX_DEL);

    gamma = P_gamma * (1.0 - P_berr * P_berr);
    gamma = MAX(gamma, MIN_VAL);
    gamma = MIN(gamma, MAX_DEL);

    Bserr = P_Bserr_1 * qd3;

    /* update cell voter ----- */
    if(Bserr < anc->voter && Bflag == FALSE) {
        output_cell = m;
        Bflag = TRUE;
    }

    Bserr = MAX(Bserr, MIN_BSERR);

    rho *= anc->lambada * SQRTF((Bserr_1 / Bserr) * (gamma / gamma_1));
    rho += berr * err;
    rho = MAX(rho, MIN_RHO);
    rho = MIN(rho, MAX_RHO);

    N_err = err - rho * berr;
}

p = (FLOAT32 *)&(anc->cells[output_cell /* *ANC_CELL_SIZE */]);
return(N_err);

```

```

VOID
SANC_Init(
    SANC_DATA    *anc)           /* input, context pointer      */
    FLOAT32      *p;
    INT32       m;

BUG1(anc);

```

```
p = (FLOAT32 *)anc->ls;
for (m = 0; m <= anc->cc; m++) {
    rho      = 0.0;
    err      = 0.0;
    ferr     = 0.0;
    berr     = 0.0;
    berr_1   = 0.0;
    delta    = 0.0;
    delta_1  = 0.0;
    Bserr    = anc->min_error;
    Bserr_1  = anc->min_error;
    gamma    = MIN_VAL;
    gamma_1  = MIN_VAL;
    p        += SANC_CELL_SIZE;
}
p = (FLOAT32 *)anc->cells;           /* Cell # 0 special case */
gamma    = 1.0;
gamma_1  = 1.0;
}
```

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant	:	Mohamed K. Diab et al.)	Group Art Unit 3305
)	
Appl. No.	:	Unknown)	
)	
Filed	:	Herewith)	
)	
For	:	SIGNAL PROCESSING APPARATUS)	
)	
Examiner	:	Unknown)	
)	

PRELIMINARY AMENDMENT
37 C.F.R. § 1.607(a)(4)

Assistant Commissioner for Patents
Washington, D.C. 20231

Dear Sir:

In the Specification:

Please amend the specification by replacing the first sentence with the following:

--This application is a continuation of U.S. Patent Application No. 08/943,511, filed October 6, 1997, which is a continuation of U.S. Patent Application Serial No. 08/572,488 (now U.S. Patent NO. 5,685,299), filed December 14, 1995, which is a continuation of Application Serial No. 08/132,812 (now U.S. Patent No. 5,490,505), filed October 6, 1993.

In the Claims:

Kindly delete Claims 1-38.

Please add the following new claim:

39. A method for measuring saturation of a blood constituent in a patient comprising the steps of:

Appl. No. : Unknown
Filed : Herewith

irradiating said patient with electromagnetic radiation of at least three discrete, different wavelengths;

sensing an intensity of said radiation for each of said wavelength after it passes through a portion of said patient to produce first, second, and third intensity signals;

representing each of said intensity signals as a function of concentration, the wavelength corresponding to the intensity signal, and a time-variable motion term corresponding to motion noise, said motion terms being proportional to one another for each of said intensity signals; and

solving the functions to obtain a value for said saturation,

wherein each of said functions includes a plurality of coefficients related to the wavelengths, the coefficients of said third function being determined based upon the coefficients of the first and second functions, and further comprising the steps of approximating at least a portion of said first and second intensity signals based upon the third intensity signal, and determining saturation from said approximation of said first and second intensity signals.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: July 7, 1998

By: Stephen C. Jensen
Stephen C. Jensen
Registration No. 35,556
Attorney of Record
620 Newport Center Drive
Newport Beach, CA 92660
(949) 760-0404

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant	:	Mohamed K. Diab, et al.)	Group Art Unit 3305
)	
Appl. No.	:	Unknown)	
)	
Filed	:	Herewith)	
)	
For	:	SIGNAL PROCESSING)	
		APPARATUS)	
)	
Examiner	:	Unknown)	
)	

REQUEST FOR INTERFERENCE
37 C.F.R. § 1.607

Assistant Commissioner for Patents
Washington, D.C. 20231

Dear Sir:

Applicant seeks to have an interference declared between this application and an unexpired patent. Pursuant to 37 C.F.R. § 1.607, Applicant submits the following information.

1. Identification of the Patent -- 37 C.F.R. § 1.607(a)(1)

The patent is U.S. Patent No. 5,645,060 ("the '060 patent") filed on June 14, 1995 and issued on July 8, 1997 to Thomas J. Yorkey, entitled METHOD AND APPARATUS FOR REMOVING ARTIFACT AND NOISE FROM PULSE OXIMETRY, and assigned to Nellcor Puritan Bennett Inc., Pleasanton, California.

2. Presentation of the Proposed Count -- 37 C.F.R. § 1.607(a)(2)

A method for measuring saturation of a blood constituent in a patient comprising the steps of:

Appl. No. : Unknown
Filed : Herewith

irradiating said patient with electromagnetic radiation of at least three discrete, different wavelengths;

sensing an intensity of said radiation for each of said wavelength after it passes through a portion of said patient to produce first, second, and third intensity signals;

representing each of said intensity signals as a function of concentration, the wavelength corresponding to the intensity signal, and a time-variable motion term corresponding to motion noise, said motion terms being proportional to one another for each of said intensity signals; and

solving the functions to obtain a value for said saturation, wherein each of said functions includes a plurality of coefficients related to the wavelengths, the coefficients of said third function being determined based upon the coefficients of the first and second functions, and further comprising the steps of approximating at least a portion of said first and second intensity signals based upon the third intensity signal, and determining saturation from said approximation of said first and second intensity signals.

An extra copy of the proposed count is submitted herewith for the Examiner's use in filling out PTO forms.

3. Identification of at Least One Claim in the Patent -- 37 C.F.R. § 1.607(a)(3)

Claims 6-10 and 15 of the '060 patent correspond to Count 1.

Appl. No. : Unknown
Filed : Herewith

4. Presentation of at Least One Claim Corresponding to the Proposed Count and An Explanation of How the Identified Claim Corresponds to the Proposed Count -- 37 C.F.R. § 1.607(a)(4)

Claim 39 of the Application corresponds to Count 1.

While Claims 6-10 and 15 of the '060 patent do not correspond exactly to the proposed count, Applicants do not currently argue that any of those claims is drawn to a separate patentable invention within the meaning of 37 C.F.R. § 1.601(n).

5. Application of the Presented Claims to the Disclosure -- 37 C.F.R. § 1.607(a)(5)

The terms of the application claims identified as corresponding to the proposed count and not previously in the application can be applied to the disclosure of the application as follows:

TERMS OF THE CLAIMS	APPLICATION TO THE DISCLOSURE
39. A method for measuring saturation of a blood constituent in a patient comprising the steps of: irradiating said patient with electromagnetic radiation of at least three discrete, different wavelengths;	Page 78, lines 19-20 FIG. 11 Page 79, lines 5-12; Page 81, lines 3-5; Page 83, lines 15-22; Page 85, lines 1-4
sensing an intensity of said radiation for each of said wavelengths after it passes through a portion of said patient to produce first, second and third intensity signals; and	Page 81, lines 3-7; Page 82, line 19 - page 84 line 1
representing each of said intensity signals as a function of concentration, the wavelength corresponding to the intensity signal, and a time-variable motion term corresponding to motion noise, said motion terms being proportional to one another; and	equations 93, 94, 95; Page 84, lines 9-34; Page 9, lines 1-26; Page 33, lines 8-23; equations 1, 2; equations 93, 94, 95; equation 25b
solving the three functions to obtain a value for said saturation;	Page 96, lines 10-31; equations 93, 94, 95

Appl. No. : Unknown
Filed : Herewith

wherein each of said functions includes a plurality of coefficients related to the wavelengths, the coefficients of said third function being determined based upon the coefficients of the first and second functions, and further comprising the steps of approximating at least a portion of said first and second intensity signals based upon the third intensity signal, and determining saturation from said approximation of said first and second intensity signals.	Page 36, lines 11-26; Page 75, line 20 - page 76, line 20; Page 77, lines 23 - 30; Equation 92; Page 84 line 8 - page 86, line 24; equations 93 - 96; Page 90, line 9 - page 93, line 22; Page 96, lines 20-31
---	---

6. Timing of the Request -- 37 C.F.R. § 1.607(a)(6)

37 C.F.R. § 1.607(a)(6) is irrelevant because this Request and the accompanying 37 C.F.R. § 1.607(a)(4) Amendment are being submitted prior to one year from the date on which the '060 patent was granted.

7. Examination Conducted With Special Dispatch -- 37 C.F.R. § 1.607(b)

Under 37 C.F.R. § 1.607(b), Applicants respectfully request that the examination of the present application be conducted with special dispatch.

8. Identification of Presented Claims Corresponding Substantially to Patent Claims - 37 C.F.R. § 1.607(c)

Applicant's Claim 39 corresponds substantially to Claim 6 of the '060 patent.

9. Request for Benefit of Filing Date of Applicant's Priority Applications

Applicant's claim priority under 35 U.S.C. § 120 to earlier filed application is as follows:

The present application is a continuation of U.S. Patent Application No. 08/943,511 filed October 6, 1997 which was a continuation of U.S. Patent Application Serial No. 08/572,488 (now U.S. Patent NO. 5,685,299) filed December 14, 1995, which is a continuation of Application Serial No. 08/132,812 (now U.S. Patent No. 5,490,505) filed October 6, 1993. Applicants are entitled to the benefit of the filing dates of these earlier filed applications for

Appl. No. : Unknown
Filed : Herewith

interference purposes because the Count reads on at least one adequately disclosed embodiment in each of the earlier applications. Indeed, the written description of the present application is identical to that of Application Serial No. 08/132,812 filed on October 6, 1993. The application of the terms of the presented claim to the disclosure is therefore equally applicable to the disclosure of Application Serial No. 08/132,812.

Applicants thus respectfully request the benefit of the priority date of October 6, 1993, the effective filing date of the present application.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: July 7, 1998

By: Stephen C. Jensen

Stephen C. Jensen
Registration No. 35,556
Attorney of Record
620 Newport Center Drive
Sixteenth Floor
Newport Beach, CA 92660
(949) 760-0404

SCJ-7458:cp/rc1/is
070698

COUNT

A method for measuring saturation of a blood constituent in a patient comprising the steps of:

irradiating said patient with electromagnetic radiation of at least three discrete, different wavelengths;

sensing an intensity of said radiation for each of said wavelength after it passes through a portion of said patient to produce first, second, and third intensity signals;

representing each of said intensity signals as a function of concentration, the wavelength corresponding to the intensity signal, and a time-variable motion term corresponding to motion noise, said motion terms being proportional to one another for each of said intensity signals; and

solving the functions to obtain a value for said saturation,
wherein each of said functions includes a plurality of coefficients related to the wavelengths, the coefficients of said third function being determined based upon the coefficients of the first and second functions, and further comprising the steps of approximating at least a portion of said first and second intensity signals based upon the third intensity signal, and determining saturation from said approximation of said first and second intensity signals.

RECORDED IN U.S. PATENT AND TRADEMARK OFFICE

FIG. 1

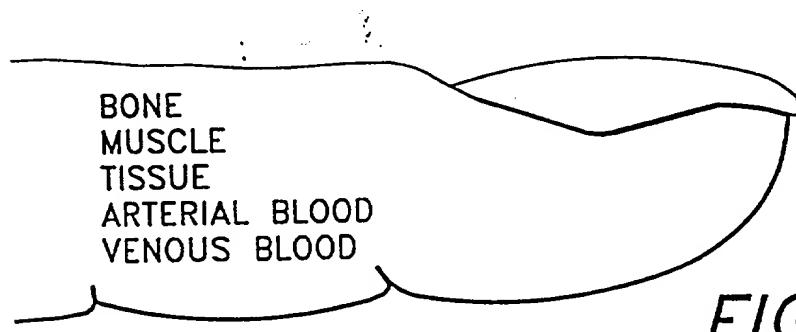
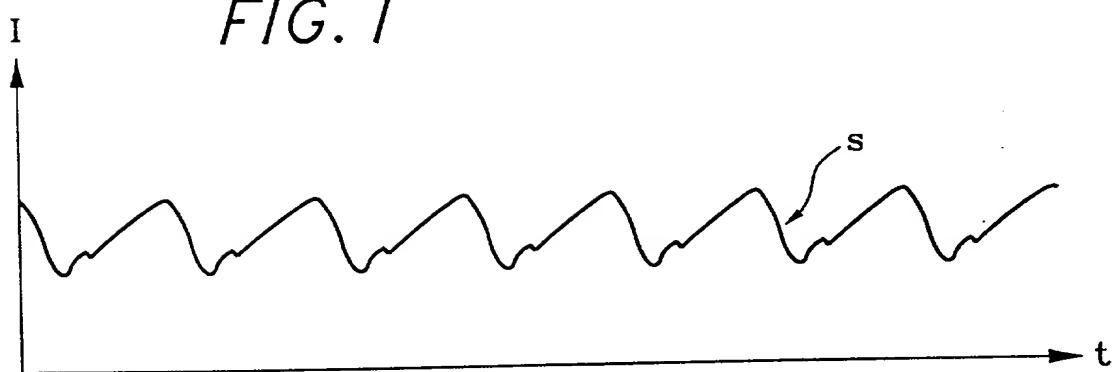
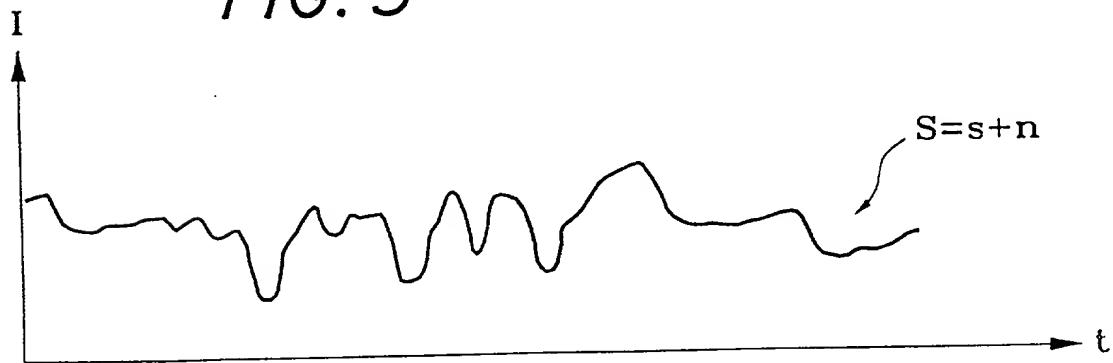


FIG. 2

FIG. 3



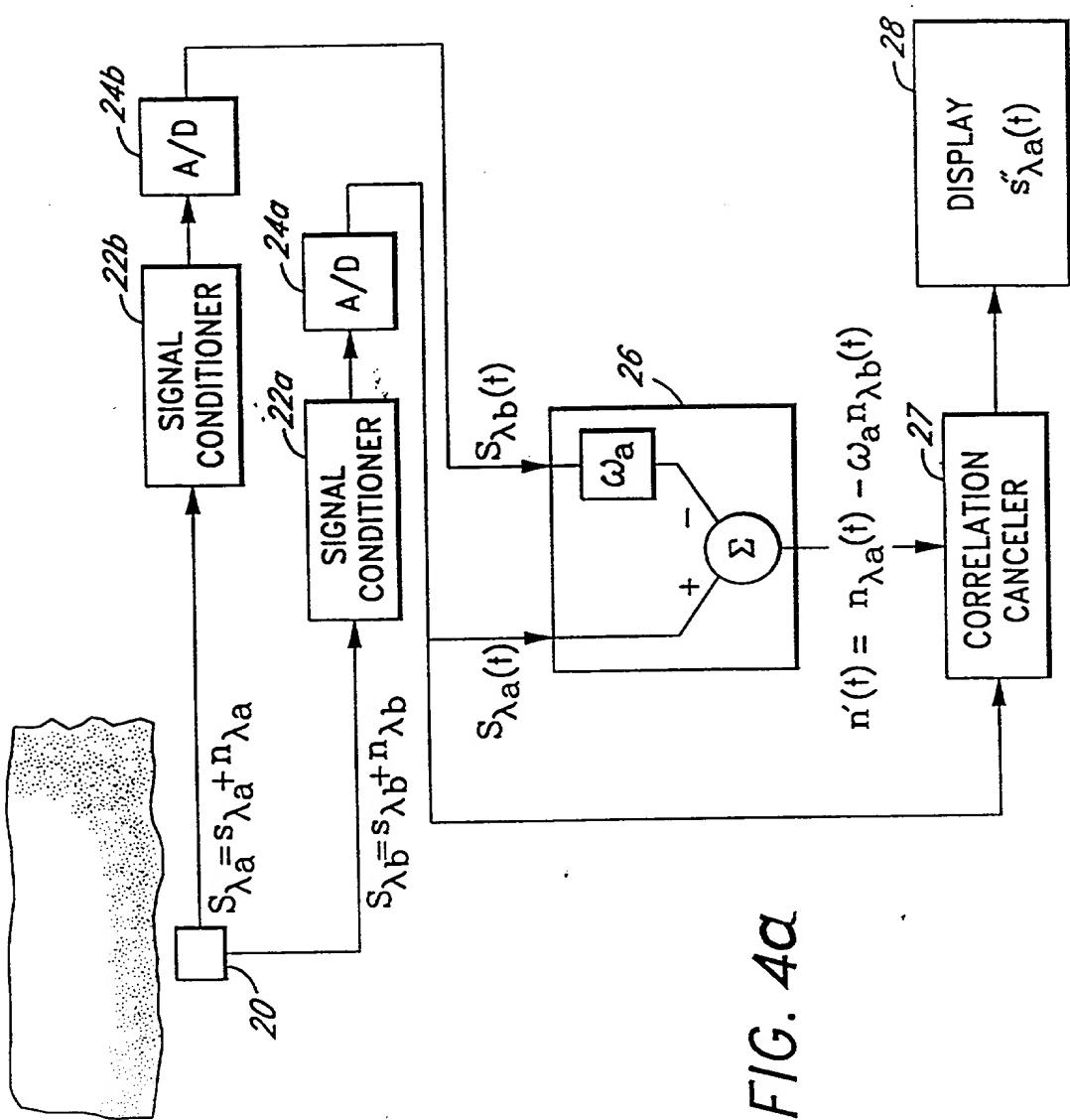


FIG. 4a

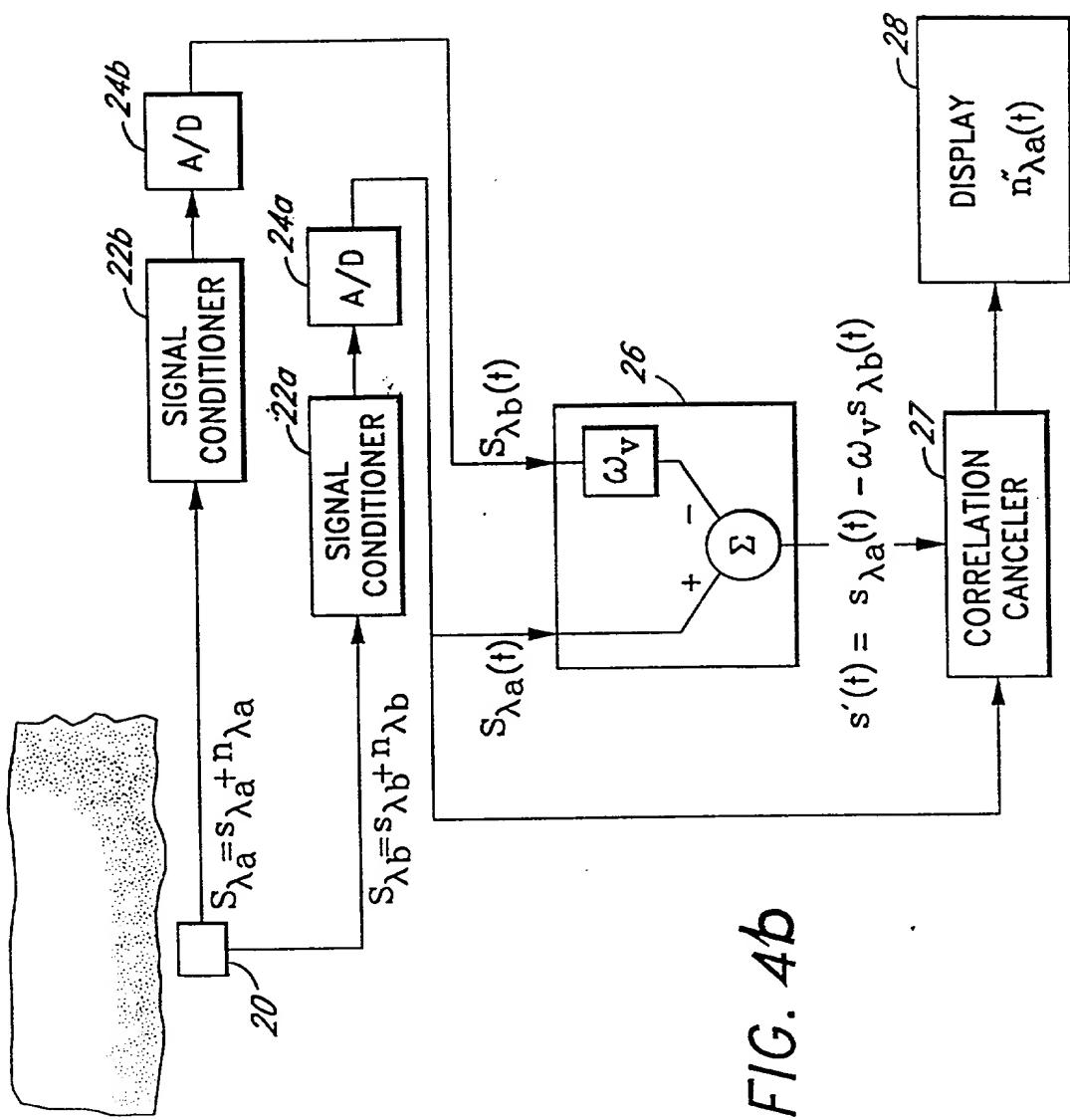


FIG. 4b

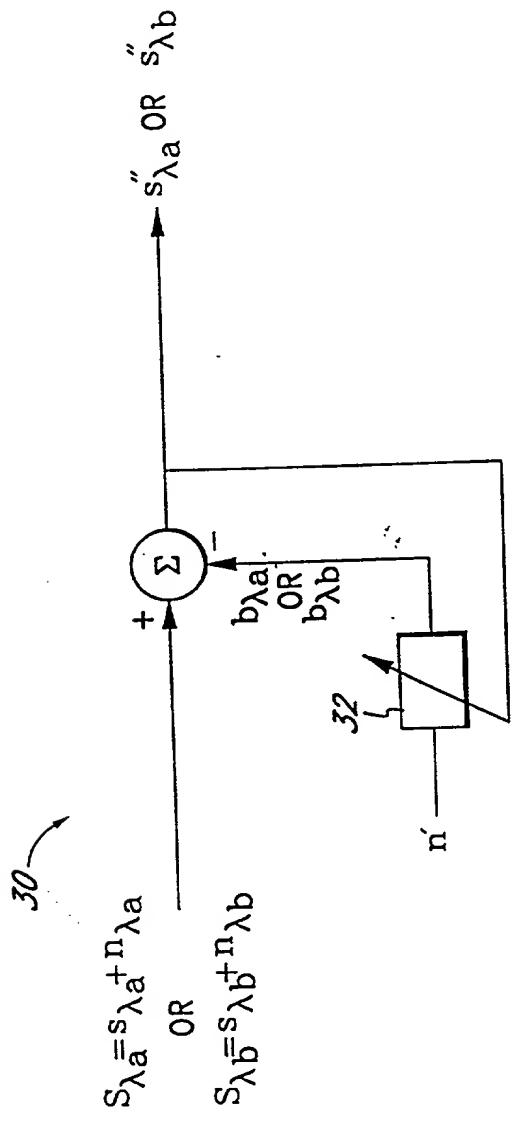


FIG. 5a

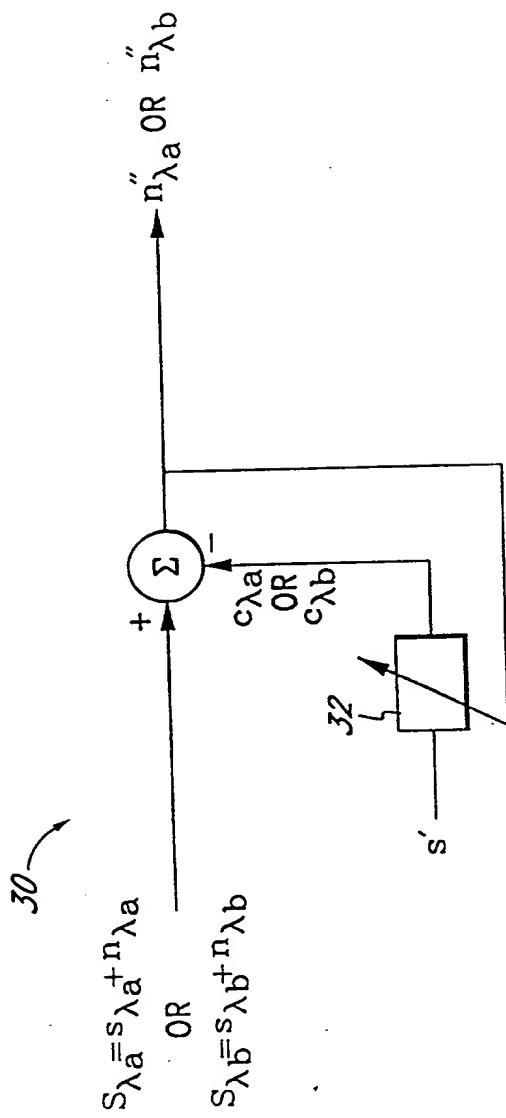


FIG. 5b

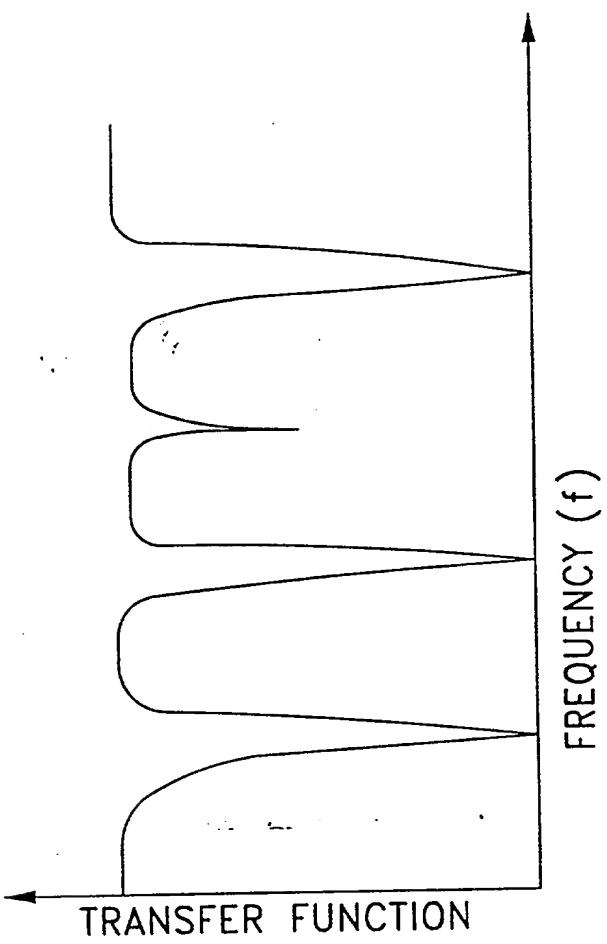


FIG. 5C

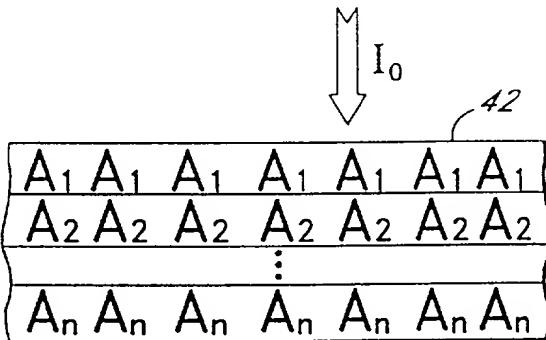


FIG. 6a

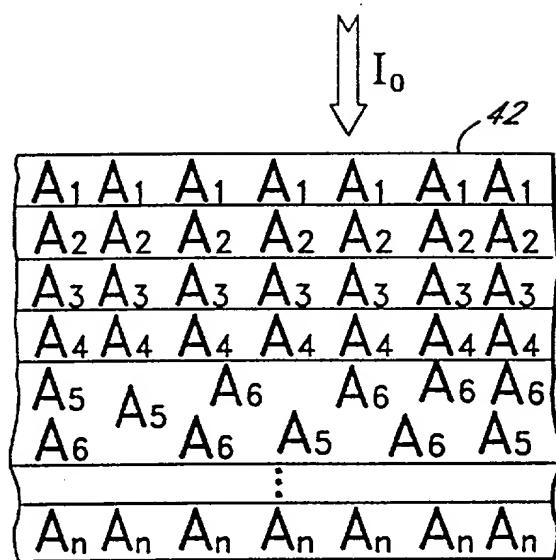


FIG. 6b

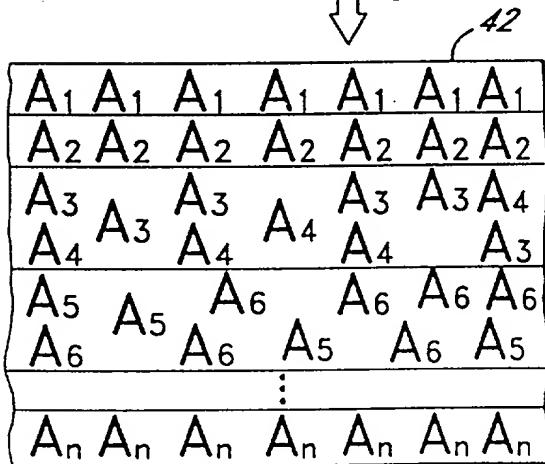
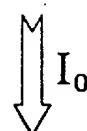
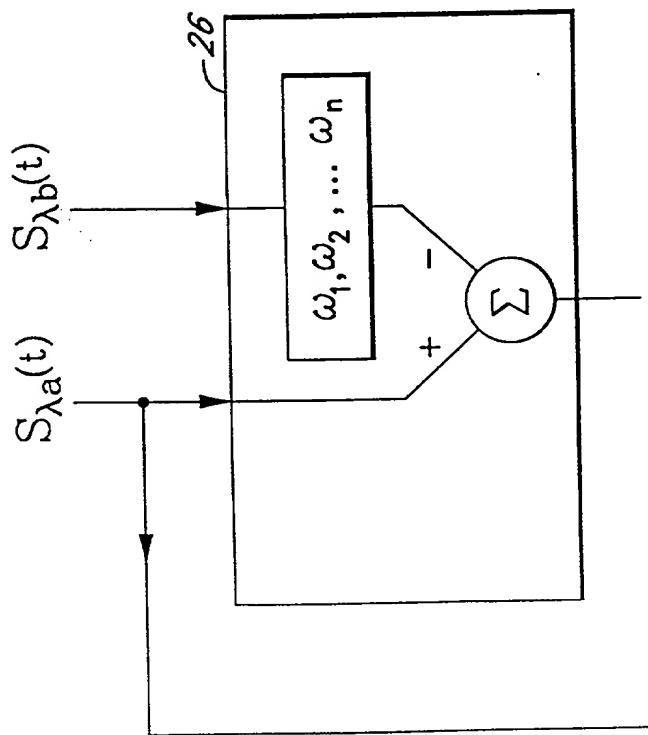


FIG. 6c



FIG. 7a



$$r'(\omega_1, t) = S_{\lambda a}(t) - \omega_1 S_{\lambda b}(t) + N_{\lambda a}(t) - \omega_1 N_{\lambda b}(t)$$

$$r'(\omega_2, t) = S_{\lambda a}(t) - \omega_2 S_{\lambda b}(t) + N_{\lambda a}(t) - \omega_2 N_{\lambda b}(t)$$

⋮

$$r'(\omega_n, t) = S_{\lambda a}(t) - \omega_n S_{\lambda b}(t) + N_{\lambda a}(t) - \omega_n N_{\lambda b}(t)$$

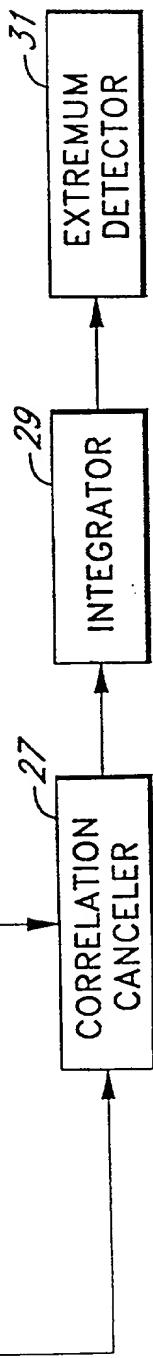


FIG. 7b

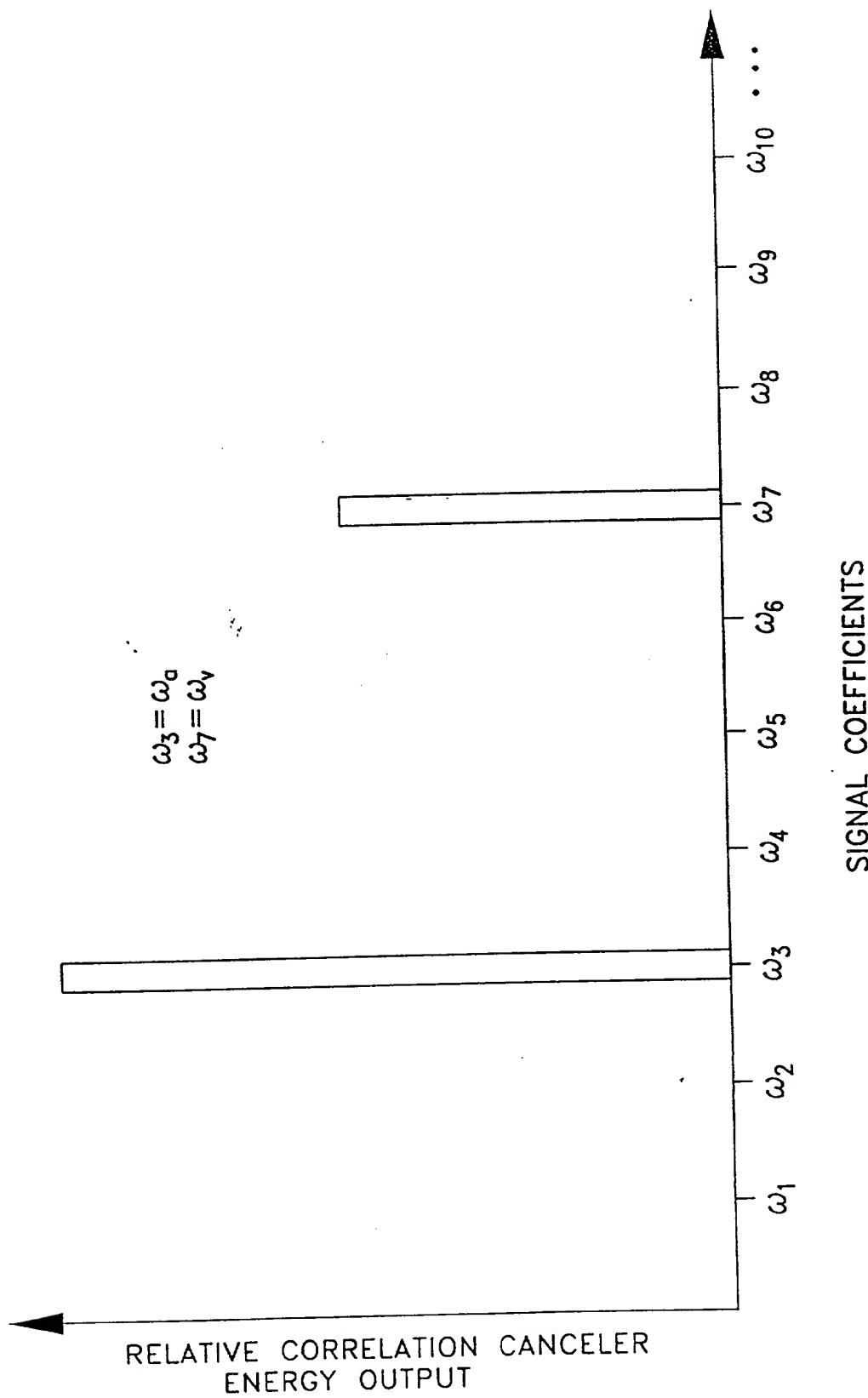


FIG. 7C

$$\begin{aligned}\omega_3 &= \omega_a \\ \omega_7 &= \omega_v\end{aligned}$$

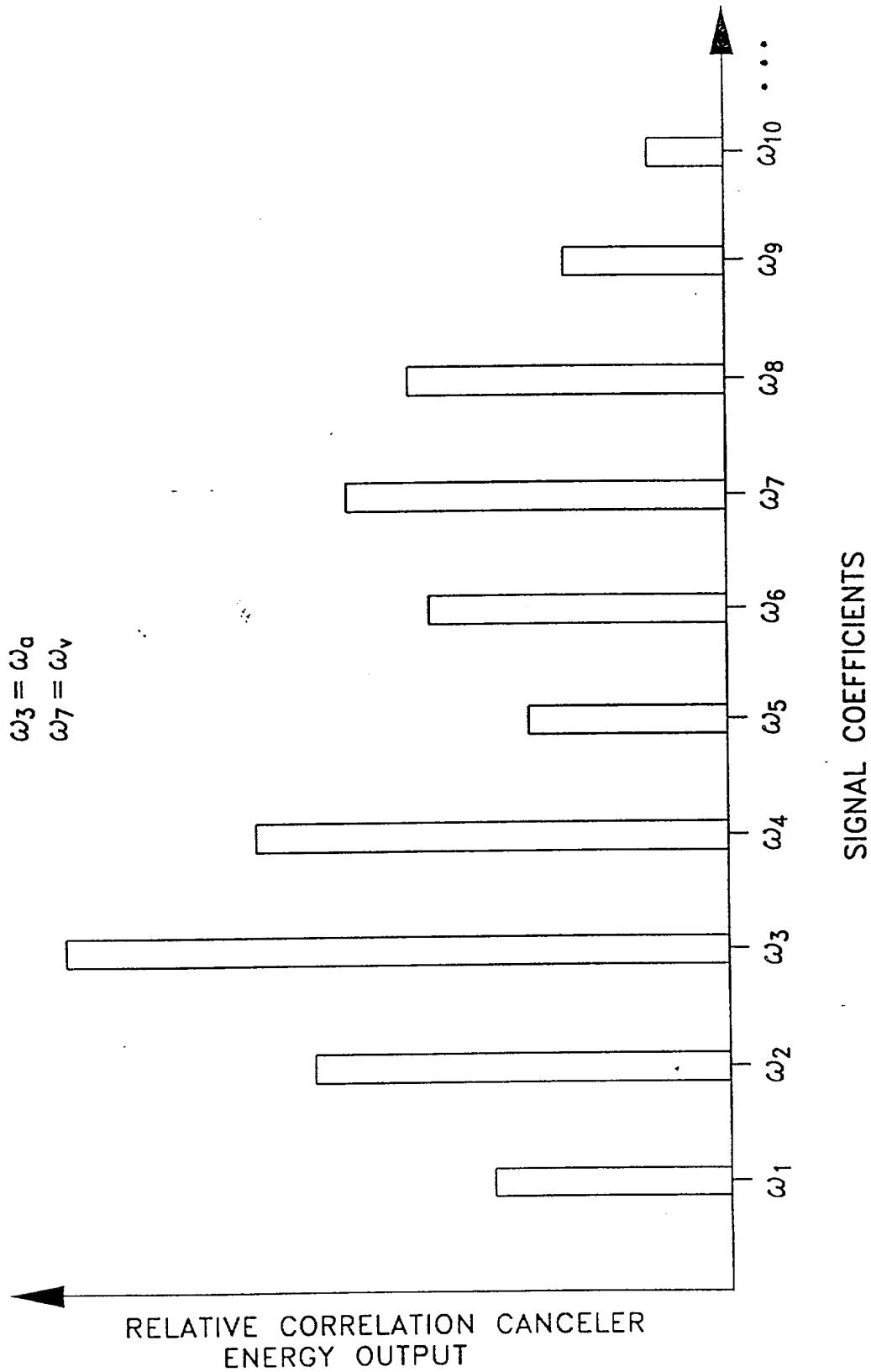
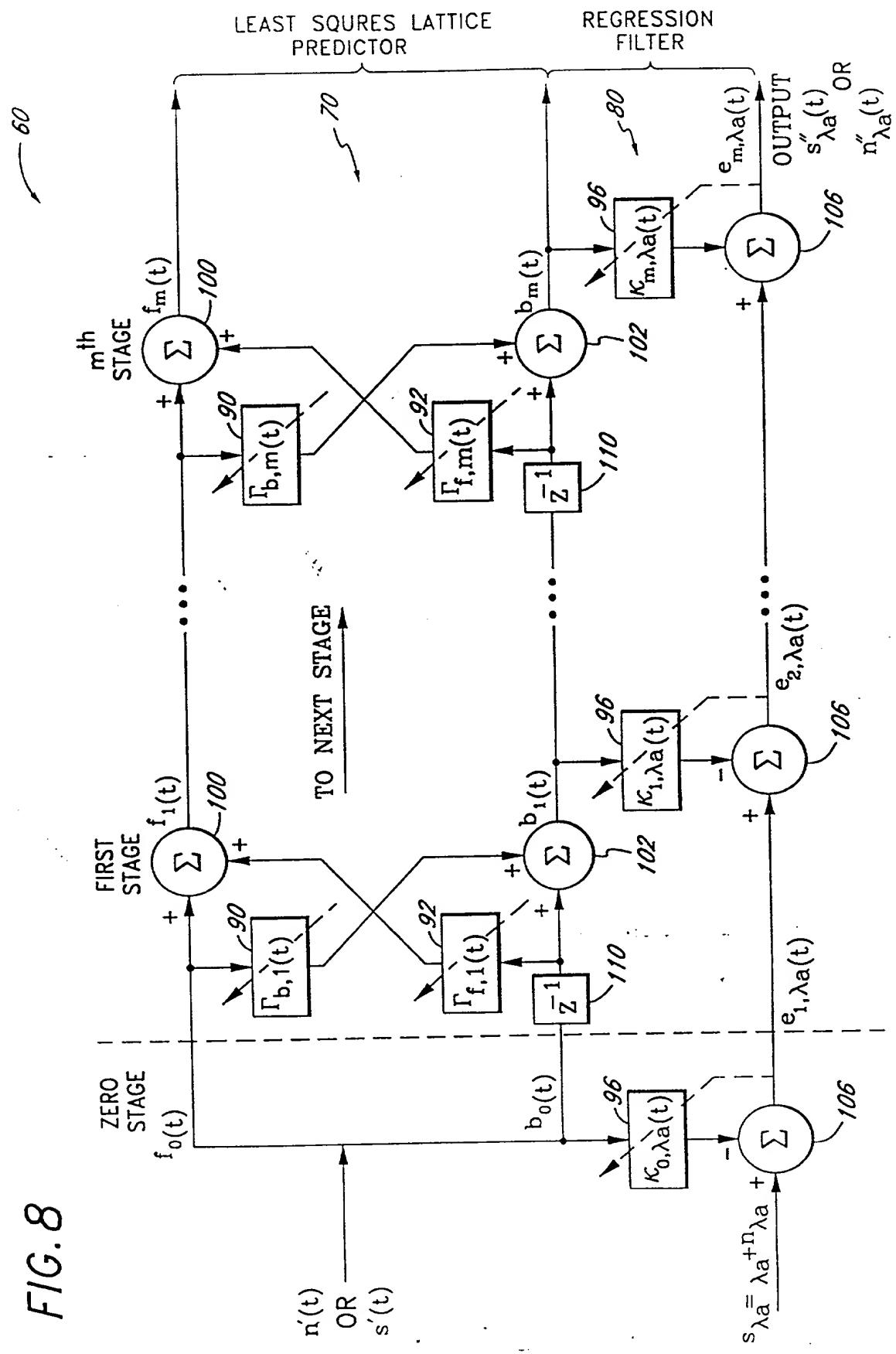


FIG. 8



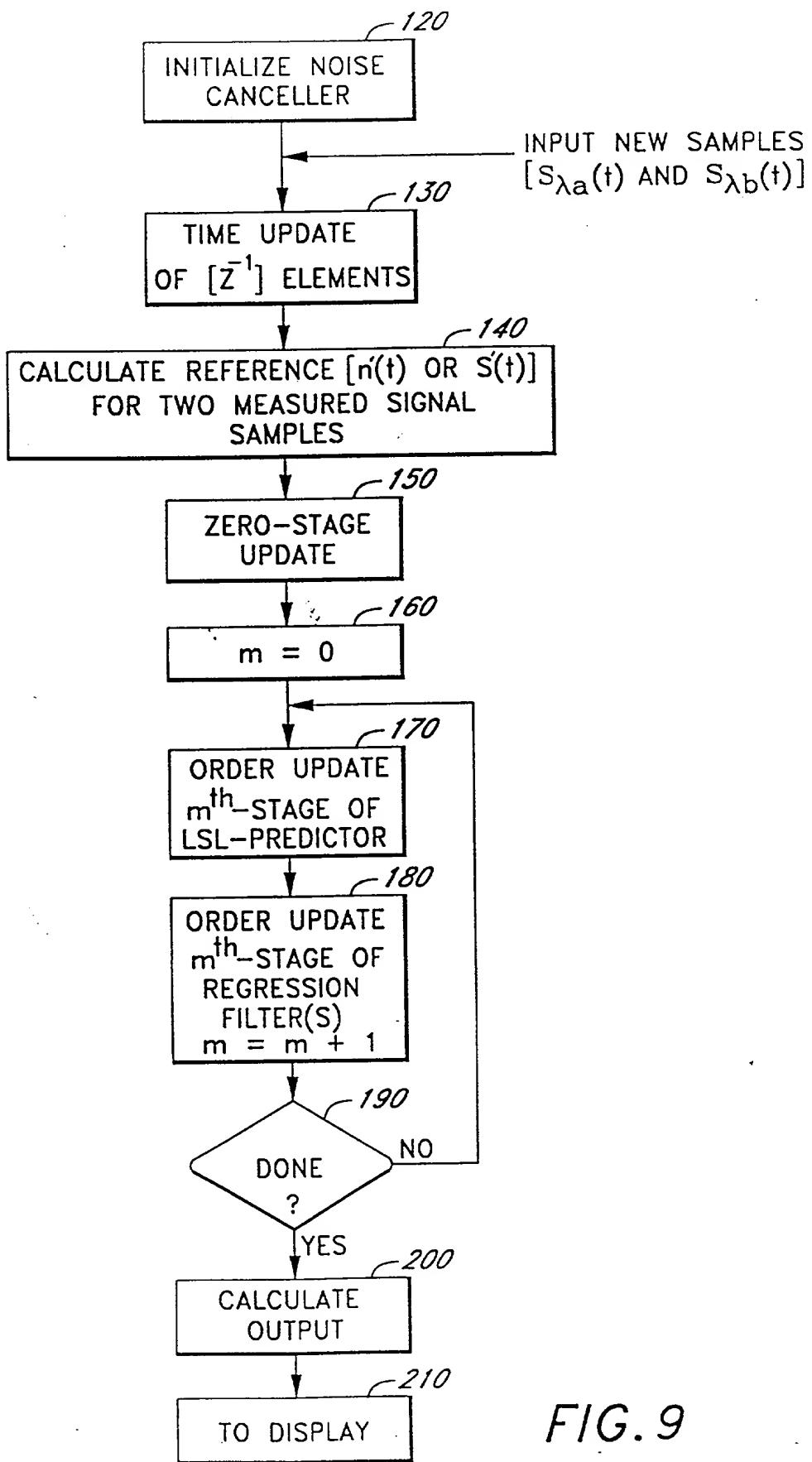
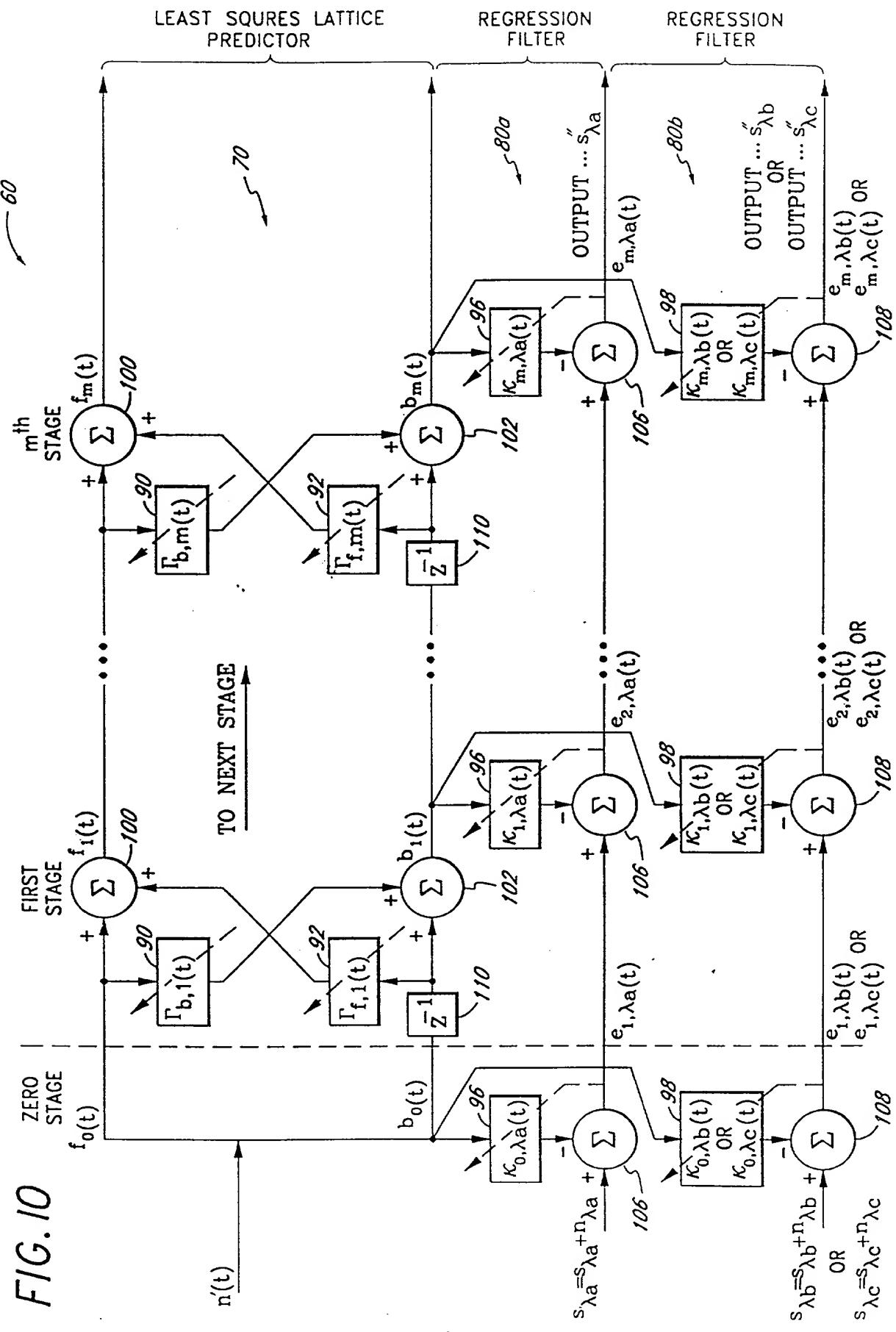


FIG. 9

FIG. 10



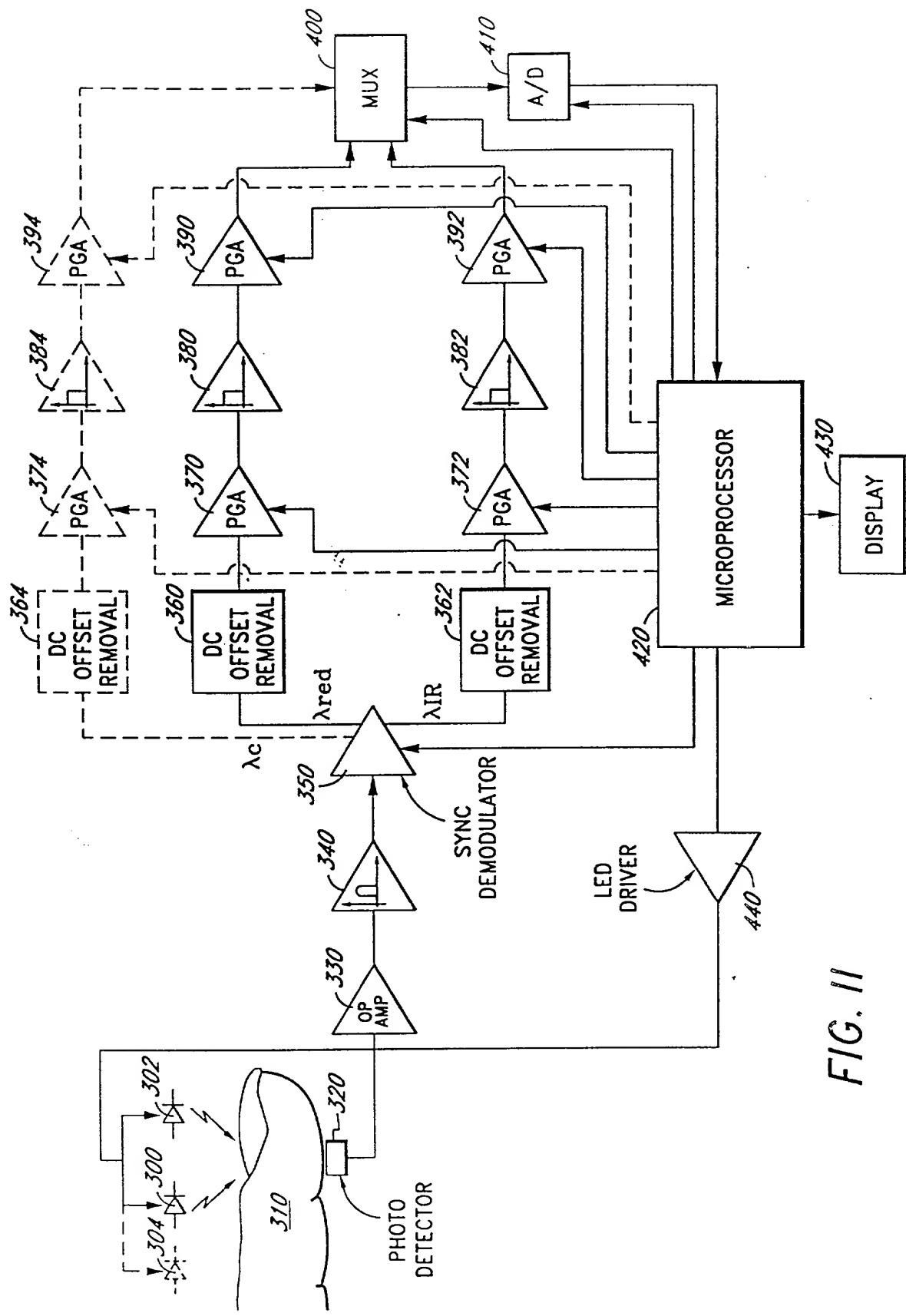
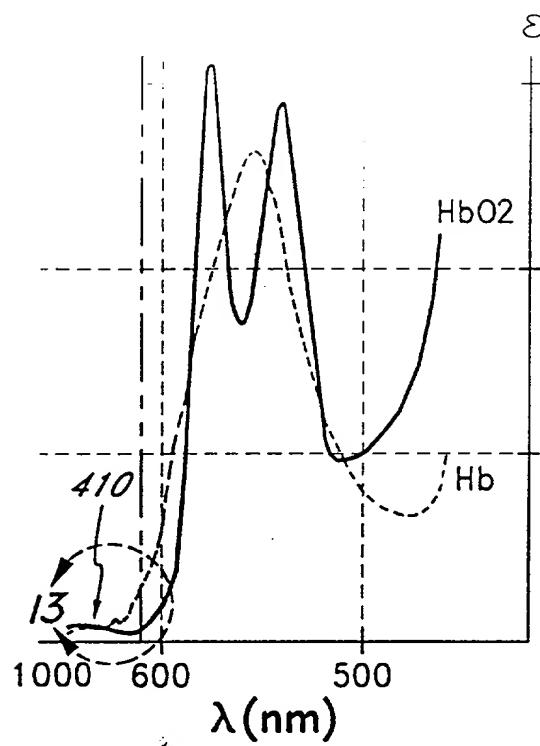


FIG. II

FIG. 12



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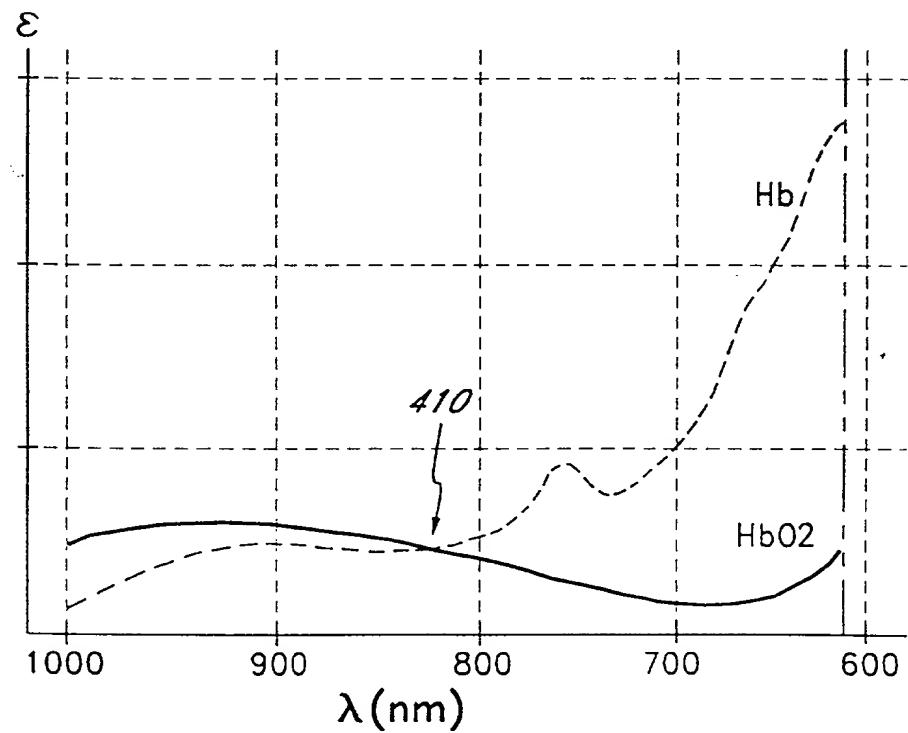


FIG. 14

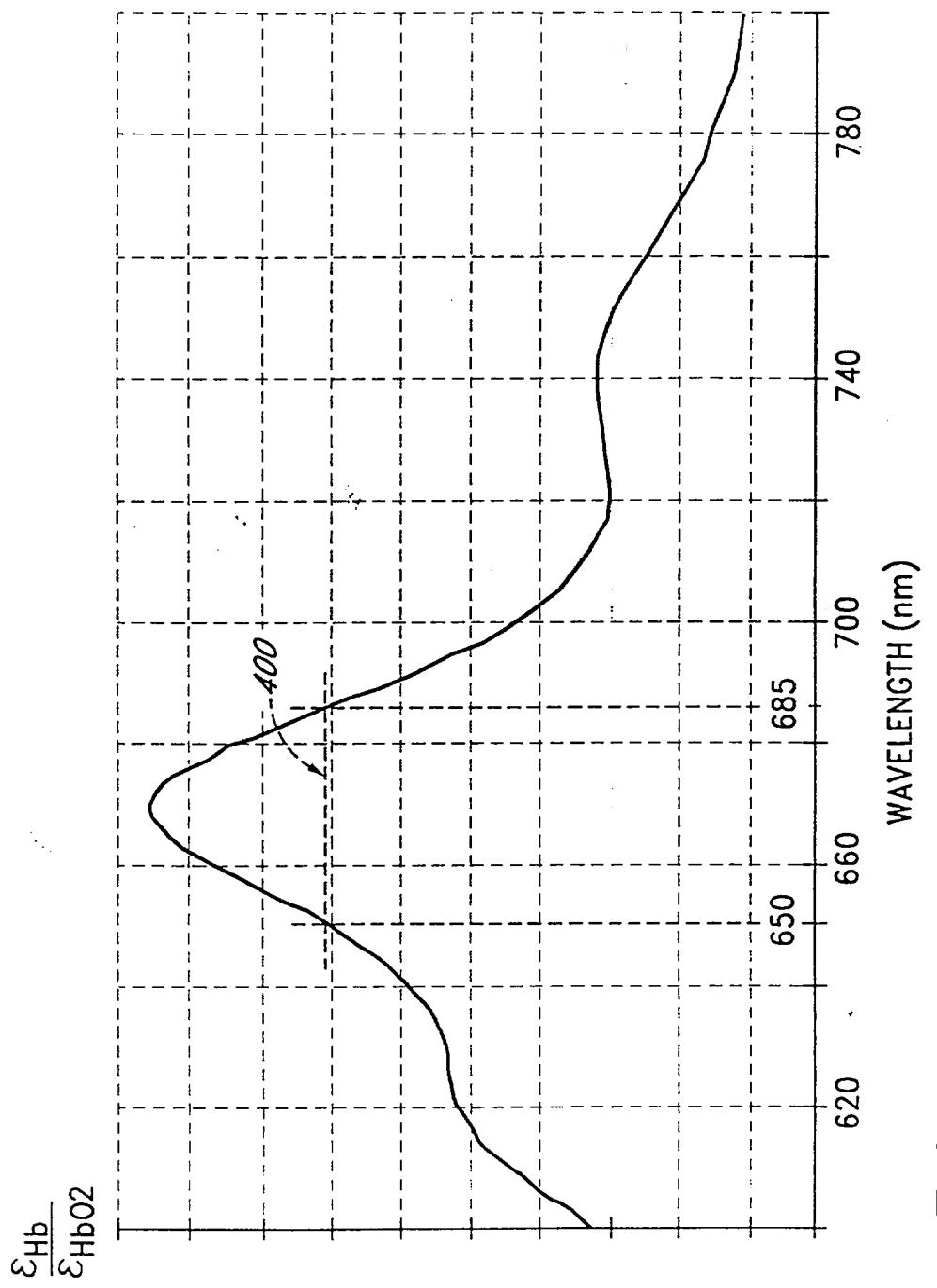
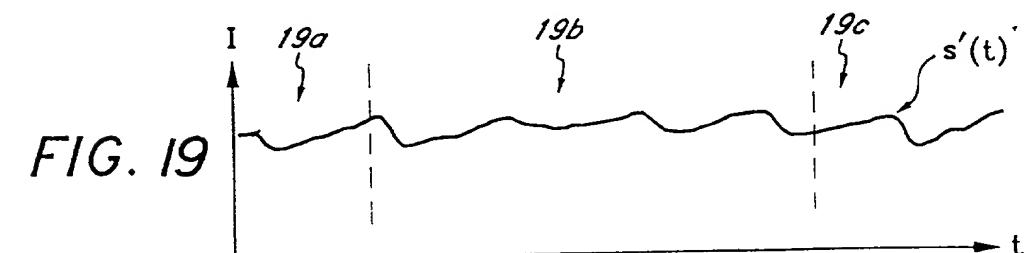
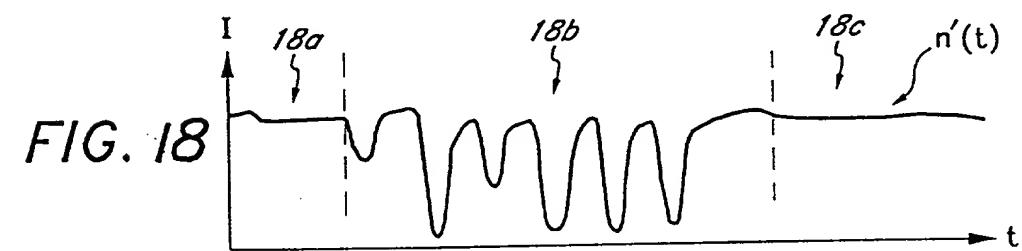
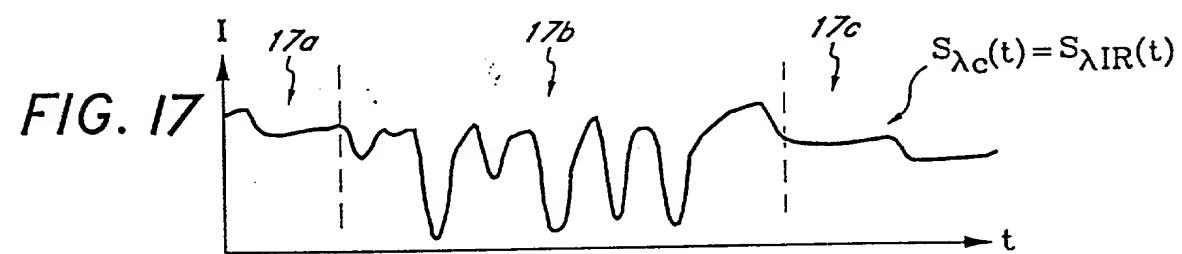
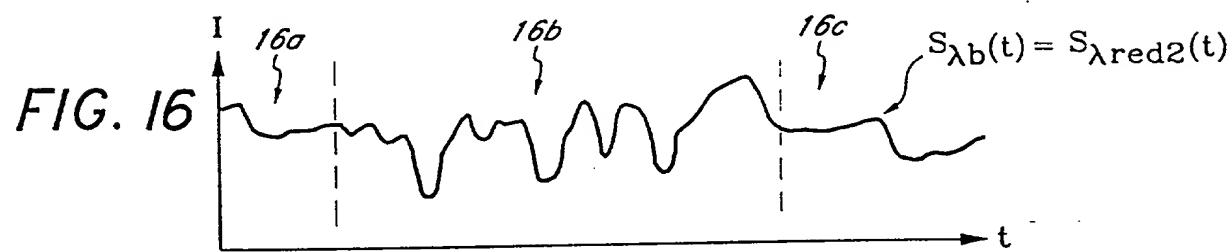
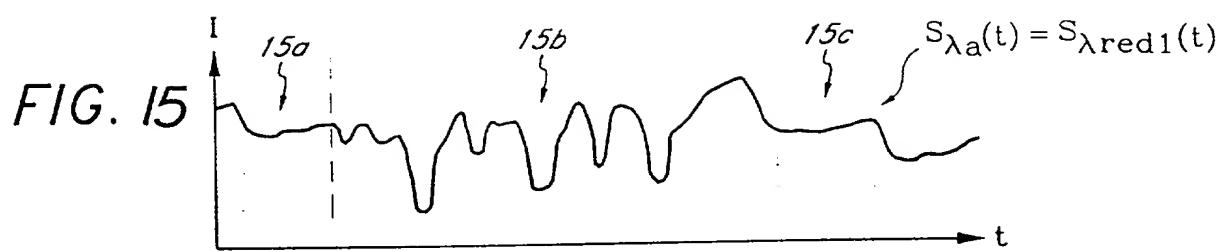


FIG. 13



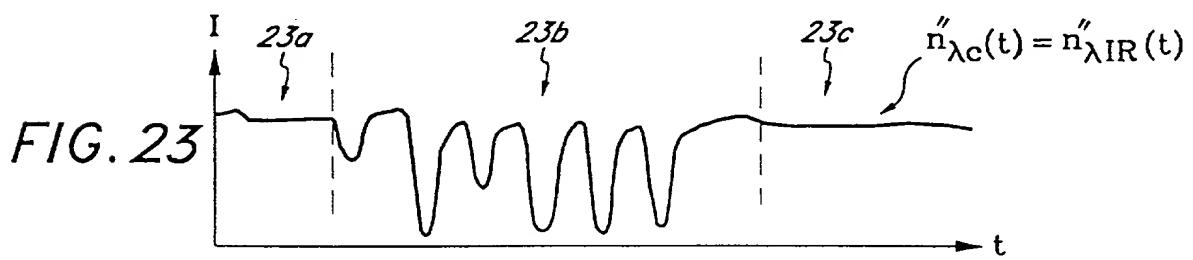
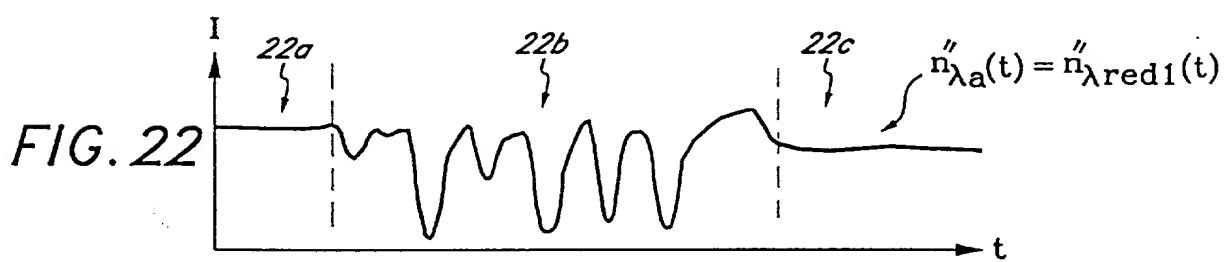
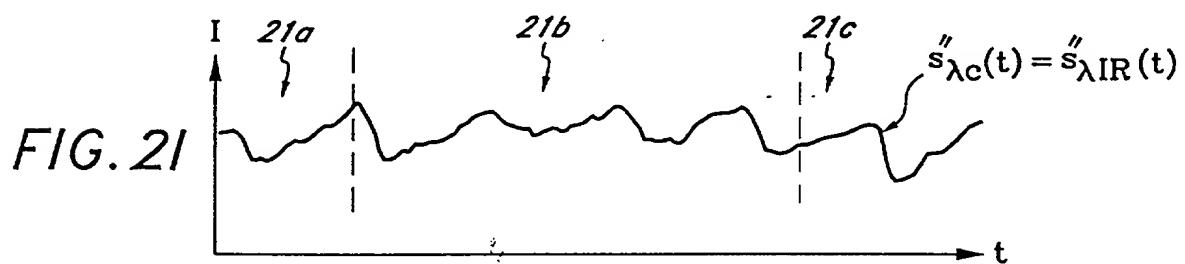
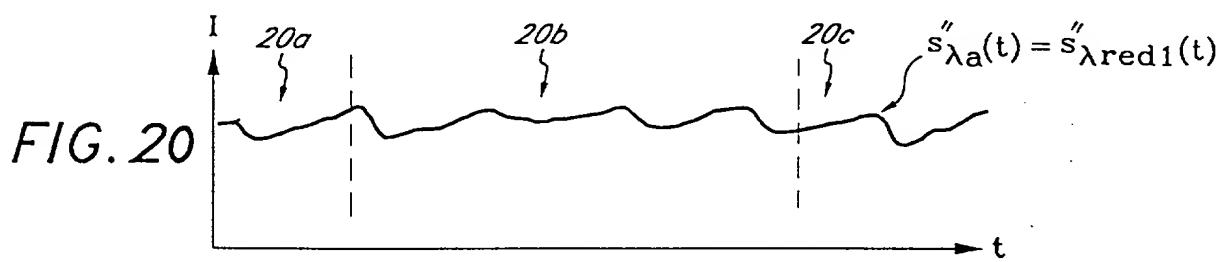


FIG. 24

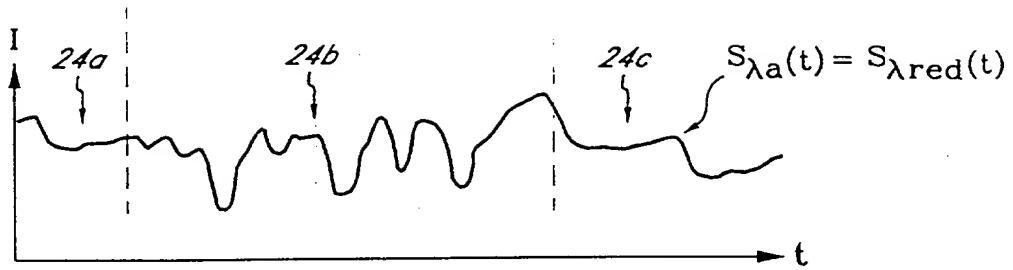


FIG. 25

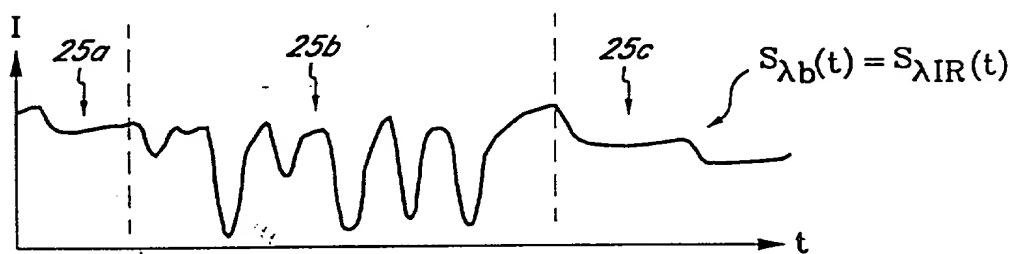


FIG. 26

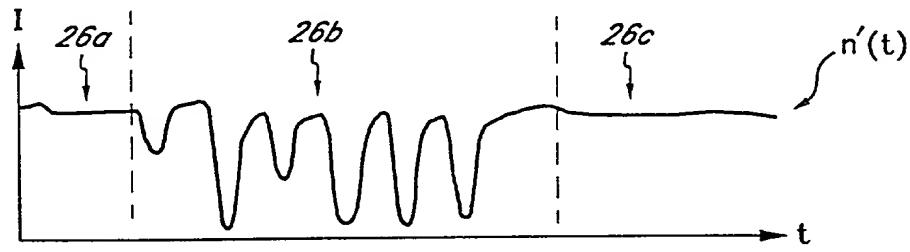


FIG. 27

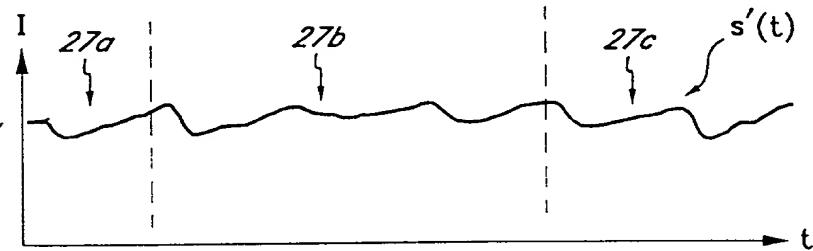


FIG. 28

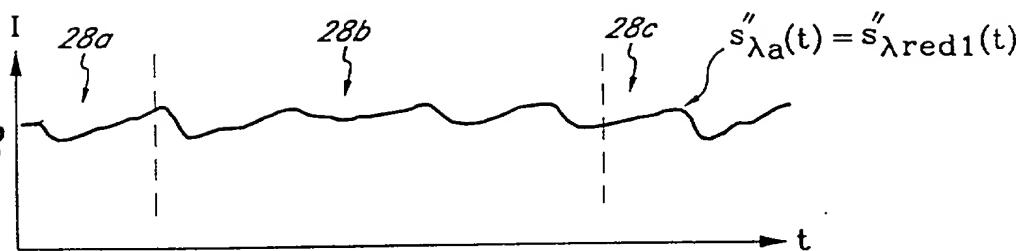


FIG. 29

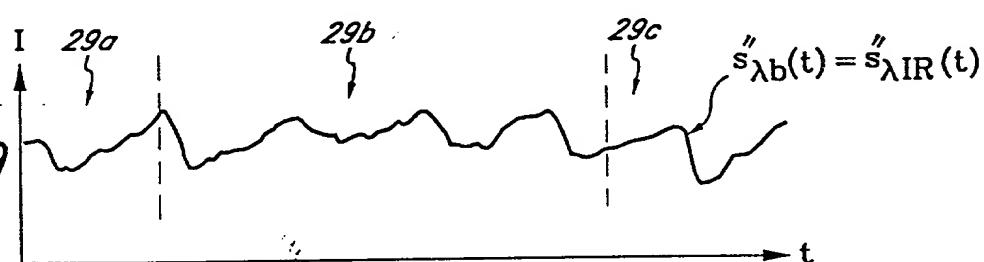


FIG. 30

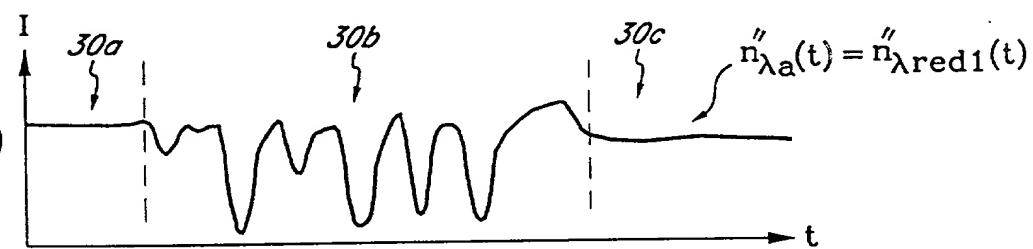


FIG. 31

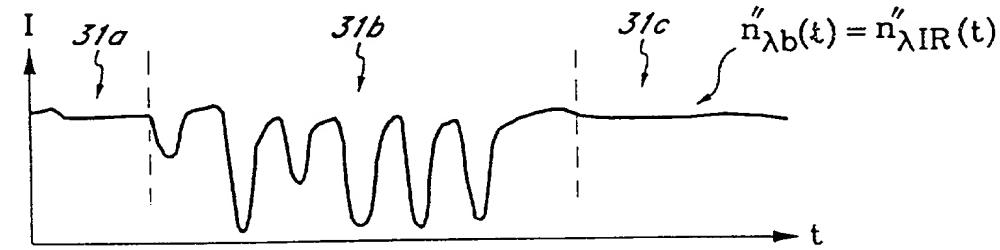
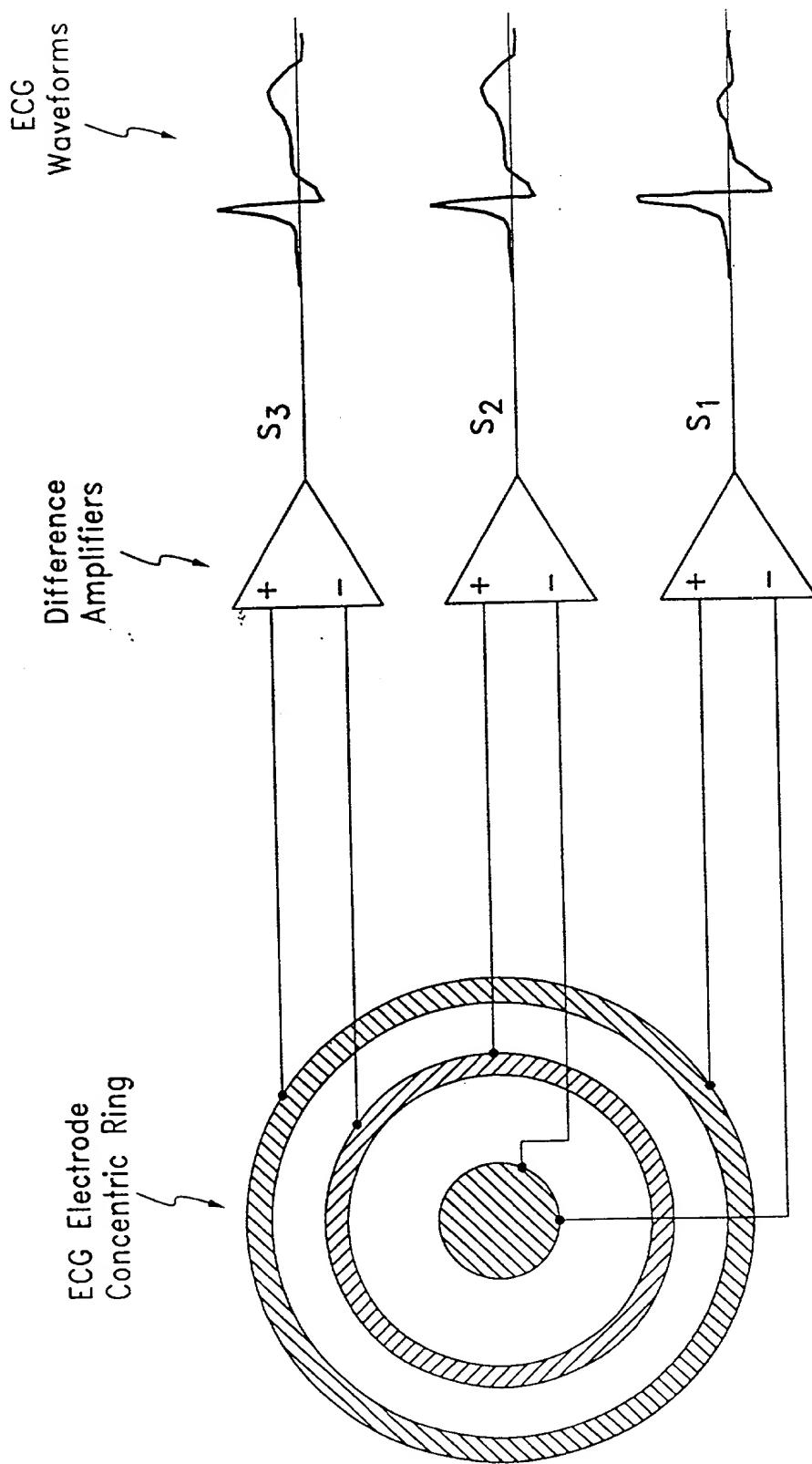


FIG. 32



COPY

PATENT
Docket No. MAS-102

DECLARATION AND POWER OF ATTORNEY

As a below named inventor of the invention claimed in the attached application for United States Letters Patent, I hereby declare that:

INVENTORSHIP IDENTIFICATION

My residence, post office address and citizenship are as stated at the end of this declaration next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the invention which is claimed in the attached application entitled:

SIGNAL PROCESSING APPARATUS

INVENTORSHIP CLAIM

I do not know and do not believe that this invention was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to this application, that this invention was not in public use or on sale in the United States of America more than one year prior to this application, that this invention has not been patented or made the subject of an inventor's certificate issued before the date of this application in any country foreign to the United States of America on an application filed by me or my legal representatives or assigns more than twelve months prior to this application, and that no application for patent or inventor's certificate on this invention has been filed in any country foreign to the United States of America prior to this application by me or my legal representatives or assigns, except as follows:

> NONE

ACKNOWLEDGEMENT OF REVIEW OF PAPERS AND DUTY OF CANDOR

I have reviewed and understand the contents of the attached application, including the specification and claims thereof, and acknowledge my duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, Section 1.56(a);

PRIORITY CLAIM

I hereby claim foreign priority benefits under Title 35, United States Code, Section 119 of any foreign application(s) for patent or inventor's certificate, or of any PCT international application(s) designating at least one country other than the United States of America, filed by me on the same subject matter as set forth in the attached application, namely:

> NONE

I hereby claim the benefits under Title 35, United States Code, section 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, section 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, section 1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

> NONE

POWER OF ATTORNEY

As a named inventor, I hereby appoint the following attorneys to prosecute this application and to transact all business in the United States Patent and Trademark Office connected therewith:

Margie R. Dickinson	Registration No. 33,679
Morland C. Fischer	Registration No. 26,188
Curtis L. Harrington	Registration No. 31,456
James E. Hawes	Registration No. 19,680

all members of the law firm of Hawes & Fischer, with offices located at 660 Newport Center Drive, Suite 460, Newport Beach, CA 92660 (tel. 714-759-6601).

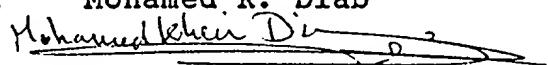
Please direct all correspondence and telephone calls to Morland C. Fischer at the above-noted address and telephone number.

DECLARATION

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

SIGNATURES

Full name of first or sole inventor: Mohamed K. Diab



Inventor's signature: _____

Date: Oct 4, 1993 Country of Citizenship: Syria

Residence: 30041 Tessier Street, #98
Laguna Niguel, California 92677

Post Office Address: 30041 Tessier Street, #98
Laguna Niguel, California 92677

Full name of second inventor: Esmaiel Kiani-Azarpayjany

Inventor's signature: 

Date: 10/4/93 Country of Citizenship: Iran

Residence: 35 Brindisi
Laguna Niguel, California 92677

Post Office Address: 35 Brindisi
Laguna Niguel, California 92677

Full name of third inventor: Walter M. Weber

Inventor's signature: 

Date: October 4, 1993 Country of Citizenship: U.S.A.

Residence: 25081 La Cresta Drive, #A
Dana Point, California 92629

Post Office Address: 25081 La Cresta Drive, #A
Dana Point, California 92629